Article
Design, Synthesis, and Characterization of Novel Coordination Compounds of Benzimidazole Derivatives with Cadmium

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Abstract: Four complexes of Cd(II) with benzimidazole derivatives were synthesized and named C1, C2, C3, and C4. All coordination compounds were characterized through elemental analysis (EA), flame atomic absorption spectrometry (FAAS), Fourier-transform infrared spectroscopy (FTIR), thermogravimetric analysis coupled with mass spectrometry (TG-MS), a cytotoxicity assay (MTT (3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazoliumbromide)), and computational chemical analysis for absorption, distribution, metabolism, and excretion (ADME). All of the obtained results are compatible and are consistent with the respective structures of the obtained compounds and their properties. The various techniques used allowed the determination of the composition, proposed structure of the compounds, their thermal stability and thermal properties, and the method of coordination between the metal (II) ion and the ligand. The ADME technique was also used to estimate the physicochemical and biological properties. The antitumor activity of the compounds was determined with an MTT assay on the glioblastoma (T98G), neuroblastoma (SK-N-AS), and lung adenocarcinoma (A549) cell lines, as well as normal human skin fibroblasts (CCD-1059Sk). Compound C2 was found to have potential antitumor properties and to be effective in inhibiting the growth of neuroblastoma cells. The antimicrobial activity of Cd complexes, free ligands, and reference drugs was tested against six strains of Gram-positive bacteria, five strains of Gram-negative rods, and three strains of yeasts. Compound C3 significantly increased activity against Gram-positive bacteria in comparison to the ligand.

Keywords: metal complexes; antitumor; antiviral; metal-based drugs; medicinal chemistry; biologically active compounds; drug discovery; pharmacological activity; lung cancer; glioblastoma; neuroblastoma; adenocarcinoma; cadmium (II) coordination compounds; benzimidazole derivatives; thermogravimetric analysis; FTIR spectra; ADME analysis; MTT assay; antimicrobial activity

1. Introduction

The creative design of new, functional coordination compounds plays a crucial role in the field of medical chemistry. An important feature of the new substances is their potential use as pharmacological agents. N-heterocyclic ligands deserve attention due to their coordination abilities and selected pharmaceutic properties. Particular attention should be paid to targeted anticancer chemotherapy. An important group of anticancer...
drugs are the coordination compounds of platinum. Their precursor and, at the same time, main representative is cisplatin. The chemotherapeutic potential of cisplatin in the treatment of cancer was first described in 1969 [1]. Currently, it is used in mono- and multi-drug therapies for the treatment of many cancers, especially testicular and ovarian cancer, head and neck cancer, and alveolar or small cell lung cancer [2–9]. Unfortunately, due to the toxic side effects (neuro- and nephrotoxicity) [10–13], narrow spectrum of action, and innate or acquired resistance of neoplastic cells to compounds of platinum [14–16], their use is limited, which is why research is carried out on chemotherapeutic agents that are based on cadmium ions together with selected organic ligands. Despite the fact that cadmium is one of the most toxic heavy metals and is in the first group of carcinogens according to the International Agency for Research on Cancer (IARC), its participation in complex compounds with anticancer activity should be considered [17–19]. The results of research on the cytotoxic properties of cadmium complexes showed that these are compounds with a significant degree of toxicity not only towards cancerous cells, but also towards normal cells; therefore, in their therapeutic effects, they resemble cisplatin [20–22]. Research on new cadmium complexes offers hope for the emergence of new and highly effective anticancer drugs, with a wide spectrum of action and a beneficial pharmacological profile. It is well known that compounds with anticancer activity that are based on endogenous metals, such as Cu and Cd, are less toxic than platinum complexes. Copper-containing coordination compounds have also been shown to be promising anticancer agents. Their activity takes various mechanisms into account, e.g., the inhibition of proteasome activity, reactive oxygen formation, telomerase activity, paraptosis, DNA intercalation, and DNA degradation [23]. This study was designed to explore the role of new derivatives as potential anticancer agents against glioblastoma (T98G), neuroblastoma (SK-N-AS), and lung adenocarcinoma (A549) cell lines, as well as normal human skin fibroblasts (CCD-1059Sk). This study confirmed that the conversion of ligands into the respective metal complexes significantly improved their anticancer properties. All of them showed much stronger activity than etoposide, which is a component of the therapy for glioblastoma, neuroblastoma, and lung cancer. There are many publications in the literature on the physicochemical and biological properties of metal coordination compounds with various derivatives of benzimidazole [24–28]. This fact proves that this is an interesting subject of research.

2. Materials and Methods

2.1. Materials and Analysis

The substrates used for ligand synthesis, as well as CdCl₂·2H₂O, were purchased from Sigma Aldrich (Warszawa, Poland). Glioblastoma (T98G), neuroblastoma (SK-N-AS), and lung adenocarcinoma (A549) cell lines and normal human skin fibroblasts (CCD-1059Sk) were obtained from the American Type Culture Collection (Manassas, VA, USA). High-glucose DMEM, DMEM/F12, MEM, fetal bovine serum, penicillin, and streptomycin were all obtained from Sigma-Aldrich.

2.2. Methods and Instruments

In order to compare the results of our work on coordination compounds with benzimidazole derivatives, the same techniques and methods were used. Samples of complexes (about 20 mg) were digested in a concentrated mixture of 36% HCl (1 mL) and 65% HNO₃ (6 mL). The contents of Cd (II) in the solid complexes were determined with an F-AAS spectrometer (Analytik Jena, contraAA 300, Jena, Germany) with a continuous source of light and using an air/acetylene flame (Analytik Jena, contraAA 300). Absorbances were measured at the analytical spectral lines at 228.8 nm for Cd (II). The limit of quantification was 0.004 mg/L for Cd (II). The solid samples were decomposed using an Anton Paar Multiwave 3000 (Graz, Austria) closed-system instrument. Mineralization was carried out for 45 min at 240 °C under a pressure of 60 bar. The contents of carbon, hydrogen, and nitrogen were determined with an instrument from Vario Micro.
Company Elementar Analysensysteme GmbH (Langenselbold, Germany). FTIR spectra were recorded with an IR Tracer-100 Schimadzu Spectrometer (4000–600 cm$^{-1}$ with an accuracy of recording of 1 cm$^{-1}$, Kyoto, Japan) using KBr pellets. The thermal analyses were carried out with a Netzsch STA 449 F1 Jupiter thermoanalyzer (Netzsch-Geratebau GmbH, Selb, Germany) coupled with a Netzsch Aeolos Quadro QMS 403 mass spectrometer (Netzsch-Geratebau GmbH, Selb, Germany). The samples were heated up to 1000 °C in corundum crucibles with a heating rate of 10 °C/min in an atmosphere of synthetic air (20% O$_2$, 80% N$_2$). The cytotoxic effects of the compounds were evaluated using glioblastoma (T98G), neuroblastoma (SK-N-AS), and lung adenocarcinoma (A549) cell lines and normal human skin fibroblasts (CCD-1059Sk). T98G and SK-N-AS cells were cultured in high-glucose DMEM and A549 cells were cultured in DMEM/F12, while CCD-1059Sk cells were cultured in MEM. All media were supplemented with 10% (FBS), 100 U/mL of penicillin, and 100 mg/mL of streptomycin (PenStrep). Cells were incubated at 37 °C in a humidified atmosphere of 5% CO$_2$. Stock solutions were prepared by dissolving the compounds in sterile dimethyl sulfoxide (DMSO) to obtain a concentration of 50 mg/mL. The cells were seeded into sterile 96-well plates (Nunc) at a density of 1 × 10$^5$ cells/mL. After 24 h of incubation, the medium was removed from each well, and then the cells were incubated for the next 24 h with different concentrations of the tested compounds (1–100 µg/mL) in their respective media containing 2% FBS. The cytotoxicity of the compounds was evaluated using an MTT assay [29], which was based on the conversion of 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) into dark-blue formazan crystals. Briefly, after 24 h of incubation of the cells with varying concentrations of the tested compounds, all culture media were removed from the plates. The cells were washed with PBS, and then 100 µL of medium containing 10% MTT solution (5 mg/mL) was added to each well. After 3 h of incubation, 100 µL (per well) of 10% SDS buffer solution was added to solubilize the formazan crystals. After overnight incubation, the absorbance was measured at 570 nm using a microplate reader (Epoch, BioTek Instruments). Experiments were repeated twice, and the measurements in each experiment were run in triplicate. The viability of the investigated cells was expressed as the percentage of the viability of the untreated cells, and the results were transformed into IC50 values (expressed as mean ± SD). The complexes were screened for antibacterial and antifungal activity by using the micro-dilution broth method with Mueller–Hinton broth for the growth of bacteria or Mueller–Hinton broth with 2% glucose for the growth of fungi [30]. The minimal inhibitory concentration (MIC) of the tested derivatives was evaluated for the panel of reference microorganisms from the American Type Culture Collection (ATCC), including Gram-negative bacteria (Escherichia coli ATCC 25922, Salmonella Typhimurium ATCC14028, Klebsiella pneumonae ATCC 13883, Pseudomonas aeruginosa ATCC 9027, Proteus mirabilis ATCC 12453), Gram-positive bacteria (Staphylococcus aureus ATCC 25923, Staphylococcus epidermidis ATCC 12228, Micrococcus luteus ATCC 10240, Enterococcus faecalis ATCC 29212, Bacillus subtilis ATCC 6633, Bacillus cereus ATCC 10876), and fungi (Candida albicans ATCC 10231, Candida parapsilosis ATCC 22019, C. glabrata ATCC 90030).

2.3. Statistical Analysis

The data were expressed as the mean ± SD values from three independent replicated experiments. P values of less than 0.05 in comparison with the control group according to Student’s t-test were considered to be statistically significant [31].

2.4. ADME Analysis

An ADME analysis was performed using the SwissADME service (Swiss Institute of Bioinformatics 2021) [32–34] and the ProTOX II service for the prediction of the toxicities of the tested compounds [35].
3. Results and Discussion
3.1. Synthesis
3.1.1. Ligand Synthesis

Ligands L1 and L2 were synthesized through the condensation and cyclization of 2,3-dimino pyridine or 3,4-diaminopyridine with isonicotinic acid in PPA (polyphosphoric acid) (Scheme 1) and ligands L3 and L4 were obtained by reacting 2,3-diaminopyridine or 5-bromo-2,3-diaminopyridine with 3-pyridinecarboxaldehyde in the presence of boric acid in a mixture of water and DMSO (Scheme 2). We previously described these syntheses [36].

\[ \text{Scheme 1. Synthesis of ligands 1 (L1) and 2 (L2).} \]

3.1.2. Complex Synthesis

The starting materials in the form of benzimidazole derivatives (0.25 mmol) and cadmium chloride dihydrates (0.25 mmol) used for the synthesis were dissolved in 96% v/v ethanol until homogeneous solutions were obtained. Then, they were mixed using a reflux condenser for about half a day. The total volume of the reaction mixture was 30 mL. The reaction was carried out under the conditions of a constant room temperature (25 °C) and controlled pH (6–7) until the formation of precipitates of the coordination compounds, which were then washed with 40% EtOH and a mixture of EtOH and H2O (vol = 1/1). The reaction products were air dried at room temperature (Scheme 3). The newly synthesized coordination compounds were characterized through an elemental C/H/N analysis, and the determination of Cd (II) content was carried out by using FTIR spectra and the TG-MS technique.

\[ \text{C1—Cd(L1)Cl}_2\cdot\text{H}_2\text{O} (\text{C}_{11}\text{H}_{10}\text{N}_4\text{O}_4\text{CdCl}_2) (397.5403 \text{ g/mol}), \text{ yield (43%}, \text{ anal. calculated (\%): Cd, 28.28; C, 33.23; H, 2.55; N, 14.09. Found (\%): Cd, 28.57; C, 33.31; H, 2.39; N, 14.25.} \]

\[ \text{C2—Cd(L2)Cl}_2\cdot\text{H}_2\text{O} (\text{C}_{11}\text{H}_{10}\text{N}_4\text{O}_4\text{CdCl}_2) (397.5403 \text{ g/mol}), \text{ yield (41%}, \text{ anal. calculated (\%): Cd, 28.28; C, 33.23; H, 2.55; N, 14.09. Found (\%): Cd, 27.88; C, 33.45; H, 2.43; N, 14.15.} \]
C3—Cd(L3)Cl2·2H2O (C11H12N4O2CdCl2) (415.5556 g/mol), yield (38%), anal. calculated (%): Cd, 27.05; C, 31.79; H, 2.91; N, 13.48. Found (%): Cd, 27.21; C, 32.00; H, 2.59; N, 13.35.

C4—Cd(L4)Cl2·2H3O (C11H11N4O2CdBrCl2) (494.4516 g/mol), yield (42%), anal. calculated (%): Cd, 22.73; C, 26.72; H, 2.24; N, 11.33. Found (%): Cd, 22.33; C, 26.37; H, 2.31; N, 11.19.

3.2. FTIR Spectra

Figures 1 and 2 show the spectra of the free ligands and the corresponding cadmium coordination compounds, respectively. During complexation, the vibrational modes of the free organic N-donors were changed. Comparing the spectra of each appropriate ligand and complex with each other, it can be seen that the fundamental $\nu$(NH) from the imidazole ring, which was present in the organic ligands in the range of 3161–2591 cm$^{-1}$ was shifted towards higher wavenumbers (3547–2928 cm$^{-1}$) in coordination compounds C2 and C4, thus showing that the NH group did not participate in binding. However, in the case of compounds C1 and C3, we can say that these bands did not change because of a lack of coordination from the imidazole ring. In the spectra of the uncoordinated donors, the vibrations modes of $\nu$(C=N) and $\nu$(C=C) were visible in the ranges of 1627–1600 and 1483–1390 cm$^{-1}$, respectively. As a result of the coordination between the metal ion and the ligand, these frequencies shifted towards higher or lower frequencies with the following peaks: for C1, 1622 and 1423 cm$^{-1}$; for C2, 1629, 1602, 1434, and 1414 cm$^{-1}$; for C3, 1598, 1450, and 1406 cm$^{-1}$; for C4, 1584, 1450, and 1395 cm$^{-1}$. For compounds C1 and C3, we could see hypsochromic effect on ligands L1 and L3, respectively. This is evidence of a different way of coordination in these coordination compounds. For all complexes, moving towards the lower wavelengths, we could observe bands in ranges of 1300–1012 and 910–690 cm$^{-1}$, corresponding to the $\beta$(CH) and $\gamma$(CH) modes, respectively. By analyzing the presented spectra, it could be concluded that, in the case of compounds C1 and C3, the coordination took place through the nitrogen atom contained in the pyridine ring and due to the slight interaction of the metal (II) and hydrogen atom coming from the imidazole ring. As for compounds C2 and C4, it was clearly seen that the N-donor ligands were monodentate and coordinated via the nitrogen from the imidazole ring.
3.3. Thermal Study

All studied compounds were hydrated; thus, the first mass loss registered on the TG curves corresponded to water removal (Figures 3 and 4a). This process finished at temperatures in the range of 180–220 °C (Table 1), which could indicate that the water molecules formed coordination bonds with the cadmium cations [37]. The dehydrated compounds started to decompose at different temperatures (Table 1), which could be a consequence of their different structures. The location of the nitrogen atoms in the two pyridine rings of C1 and C2 favored the formation of polymeric compounds, whereas in C3 and C4, it rather limited the propagation of the infinite structure. The thermal stability of coordination polymers is usually greater than that of discrete coordination compounds created from similar ligands [38]; hence, the structural diversity could be the most probable explanation for the greater thermal stability of C1 and C2 in comparison to that of C3 and C4. The decomposition pathway of the organic ligand in the studied cadmium compounds was similar to that reported for analogous copper compounds [36]. However, the initial decomposition temperatures are generally higher for cadmium compounds. Firstly, the non-conjugated pyridine ring decomposed. The mass spectra at this stage for all of the studied compounds contained signals that were characteristic of pyridine fragmentation ions, i.e., HCN⁺ and C₄H₄⁺ [39], as well as of the combustion of heterocyclic organic moieties, i.e., H₂O⁺, NO⁺, and CO₂⁺ (Figure 4b). At temperatures in the range of 460–475 °C, the second
step of organic ligand decomposition began (Table 1). For only C1, it was not possible to establish borders between the ligand decomposition substages due to their overlapping (Figure 3). The second substage of ligand decomposition was accompanied by the partial removal of the cadmium content in the form of cadmium chloride, a random amount of which evaporated above its melting point (564°C) [40]. Therefore, the residual mass of cadmium oxide as a final product was lower than expected for C1–C3 (Table 1). The calculated mass of cadmium oxide was above 30%. The presence of signals of Cl⁺ and Cl₂⁺ ions in the mass spectra in this stage (Figure 4c) was evidence of the evaporation of CdCl₂, which most likely underwent disintegration in the mass spectrometer. In the case of C4, total decomposition occurred. The entire cadmium content was removed in the form of both CdCl₂ and CdBr₂ (melting point: 568°C) [40]. Bromide ions were generated during the decomposition of the imidazopyridine part of the ligand. Similarly to CdCl₂, the signals of Br⁺ and Br₂⁺ derived from CdBr₂ were present in the mass spectra of the last decomposition stage of C4 (Figure 4d). The total decomposition of cadmium coordination compounds containing bromine is a phenomenon that is well known in the literature [41].

Figure 3. Cont.
Figure 3. Thermal analyses of the cadmium (II) coordination compounds. TG (green), DTA (blue), and DTG (red) curves for complexes C1, C2, C3, and C4.

Figure 4. Cont.
Figure 4. Representative mass spectra of volatile products of the first (a) and the second (b) stages of the thermal decomposition of C1–C4, as well of the third stage of the thermal decomposition of C1–C3 (c) and C4 (d). The correction of the mass spectra included subtracting the background spectrum and the application of automatic software correction for the carrier gas.
Table 1. Temperature ranges and mass losses of the thermal decomposition stages of the cadmium (II) coordination compounds.

|          | C1             | C2             | C3             | C4             |
|----------|----------------|----------------|----------------|----------------|
| Stage I  | 35–220 °C      | 35–220 °C      | 35–180 °C      | 35–185 °C      |
|          | 3.7%           | 4.1%           | 7.3%           | 8.6%           |
| Stage II | 430–760 °C     | 360–475 °C     | 265–465 °C     | 320–453 °C     |
|          | 88.8%          | 26.8%          | 27.6%          | 14.6%          |
| Stage III| 475–705 °C     | 465–715 °C     | 460–775 °C     |                |
|          | 58.5%          | 60.2%          | 76.8%          |                |
| Residual | 7.5%           | 10.6%          | 4.9%           |                |

3.4. MTT Cytotoxicity Assay

The ligands alone (L1, L2, L3, L4) and their Cd (II) metal complexes were screened for anticancer activity against the glioblastoma, neuroblastoma, and lung carcinoma cell lines (T98G, SK-N-AS, A549, respectively). Moreover, in order to assess the selectivity of the anticancer effect, the investigated compounds were also evaluated against normal human skin fibroblasts (CCD-1059Sk). Etoposide, which is a topoisomerase IIα inhibitor, was used as a reference drug. The results obtained confirmed that the conversion of the ligands into the respective metal complexes significantly improved their anticancer properties (Table 2). All of them showed much stronger activity than that of etoposide, which is a component of the therapy for glioblastoma, neuroblastoma, and lung cancer. Unfortunately, increased cytotoxicity of the metal complexes was also observed in relation to normal cells. Importantly, the neuroblastoma SK-N-AS cells, unlike T98G and A549 cells, exhibited greater sensitivity to the investigated cadmium complexes when compared to normal human cells. The most promising Cd (II) complex, i.e., C2, inhibited the growth of neuroblastoma cells five times more than etoposide, with an IC_{50} equal to 22.46 µM (vs. 115.25 µM for etoposide).

Table 2. Cytotoxic effects of the investigated compounds against glioblastoma (T98G), neuroblastoma, and lung carcinoma cell lines and normal human fibroblasts (CCD-1059Sk) determined with an MTT assay after 24 h of incubation.

| Complex | IC_{50} ± SD [µM] |
|---------|-------------------|
| L1      | 210 ± 12          |
| C1      | Not determined due to low solubility in culture medium |
| L2      | 178.3 ± 7.3       |
|         | 415 ± 17          |
|         | 220 ± 11          |
| C2      | 65.25 ± 0.63      |
|         | 22.5 ± 2.3        |
|         | 34.13 ± 0.55      |
|         | 44.07 ± 0.33      |
| L3      | >509              |
|         | >509              |
|         | >509              |
| C3      | 55.7 ± 2.5        |
|         | 34.53 ± 0.41      |
|         | 47.2 ± 2.1        |
|         | 40.38 ± 0.41      |
| L4      | >362              |
|         | >362              |
|         | >362              |
| C4      | 42.53 ± 0.96      |
|         | 25.3 ± 1.0        |
|         | 37.3 ± 2.1        |
|         | 27.66 ± 0.90      |
| Etoposide | >170             |
|         | 115.3 ± 3.5       |
|         | >170              |
|         | >170              |

3.5. Antimicrobial Activity

The antimicrobial activity of the Cd complexes, free ligands, and reference drugs was tested against six strains of Gram-positive bacteria, five strains of Gram-negative rods, and three strains of yeasts. The antimicrobial properties were expressed as the MIC (minimum inhibitory concentration) in mg/L (Table 3). The antimicrobial activity of the Cd complexes was compared with the antimicrobial and antifungal properties of the appropriate ligands. Vancomycin (Van), ciprofloxacin (Cip), and nystatin (Nys) were used as the standard drugs. The tested ligands and complexes showed no bioactivity against Gram-negative bacteria, except for C3, which had significantly increased activity in comparison to its ligand. Against
Gram-positive bacteria, moderate bioactivity was detected, with only a slight increase for staphylococci. Thus, the antibacterial efficiency of the tested complexes against Gram-positive bacteria decreased in the following order: $C_1 > C_2 > C_4 > C_3$. Notably, all tested Cd complexes revealed good activity against yeasts in comparison to their ligands (MIC in the range of 7.8–125 mg/L).

### Table 3. In vitro antimicrobial activity of the tested ligands and complexes.

| Microorganism       | Chemicals | L1  | C1  | L2  | C2  | L3  | C3  | L4  | C4  | Ref Drugs |
|---------------------|-----------|-----|-----|-----|-----|-----|-----|-----|-----|-----------|
| Gram-positive bacteria | Van       | >1000 | 500 | 1000 | 1000 | >1000 | 1000 | >1000 | 1000 | 0.98      |
| S. aureus ATCC 25923  |           | >1000 | 250 | 1000 | 500 | >1000 | 1000 | 1000 | 250 | 0.98      |
| S. epidermidis ATCC 12228 |         | 500  | 7.8 | 250  | 7.8 | 500  | 7.8 | >1000 | 7.8 | 0.12      |
| M. luteus ATCC 10240  |           | >1000 | 125 | 1000 | 125 | >1000 | 1000 | 1000 | 250 | 1.95      |
| E. faecalis ATCC 29212 |         | >1000 | 7.8 | 500  | 0.98 | 1000 | 7.8 | 1000 | 7.8 | 0.24      |
| B. subtilis ATCC 6633  |           | >1000 | 500 | 250  | 500 | >1000 | 1000 | 1000 | 250 | 0.98      |
| B. cereus ATCC 10876  |           | >1000 | 7.8 | 500  | 0.98 | 1000 | 7.8 | 1000 | 7.8 | 0.24      |
| Gram-negative bacteria | Cip       | >1000 | 1000 | 1000 | >1000 | 125 | >1000 | 1000 | 0.061     |
| S. typhimurium ATCC 14028 | | >1000 | 1000 | 1000 | >1000 | 125 | >1000 | 500  | 0.015     |
| E. coli ATCC 25922    |           | >1000 | 1000 | 1000 | >1000 | 31.3 | >1000 | 1000 | 0.03      |
| P. mirabilis ATCC 12453 |         | >1000 | 1000 | 1000 | >1000 | 1000 | >1000 | 1000 | 0.12      |
| K. pneumoniae ATCC 13883 |       | >1000 | 1000 | >1000 | >1000 | 250 | >1000 | 1000 | 0.49      |
| P. aeruginosa ATCC 9027 |         | >1000 | >1000 | >1000 | >1000 | >1000 | >1000 | >1000 | >1000 |          |
| Yeasts               | Nys       | >1000 | 62.5 | >1000 | 31.3 | >1000 | 31.3 | >1000 | 31.3 | 0.48      |
| C. glabrata ATCC 2091 |           | >1000 | 500 | 1000 | 125 | >1000 | 125 | 500  | 125 | 0.24      |
| C. albicans ATCC 102231 |         | >1000 | 15.6 | >1000 | 7.8 | 1000  | 15.6 | 1000 | 7.8 | 0.24      |
| C. parapsilosis ATCC 22019 |      | >1000 | >1000 | >1000 | >1000 | >1000 | >1000 | >1000 | >1000 |          |

3.6. ADME Analysis

Bioavailability radars were used for all complexes (Figure 5). The pink-colored zone on the bioavailability radar was made in the SwissADME service; it represents the optimal range for each property to indicate the drug-likeness of a molecule: Lipophilicity (LIPO) was within the range $-0.7 < \text{XlogP3} < +5.0$; molecular weight (SIZE) was within the range $150 \text{ g/mol} < \text{MW} < 500 \text{ g/mol}$; polarity (POLAR) was within the range $20 \text{ Å}^2 < \text{TPSA} < 130 \text{ Å}^2$; insolubility (INSOLU) was within the range $0 < \text{logS} < 6$; insaturation (INSATU) was within the range $0.25 < \text{Fraction Csp3} < 1$; flexibility (FLEX) was within the range $0 < \text{Num. rotatable bonds} < 9$. The non-dimer complexes met the rules of Lipinski [42], Ghose [43], Egan [44], Veber [45], and Muegge [46]. Only the dimer complex $C_3$ did not indicate good brain penetration or oral bioavailability. Coordination compound $C_3$ was too large, and its molecular weight exceeded the limit of 800 g/mol (831.11 g/mol) for being an obvious candidate as a drug. Gastrointestinal absorption and brain access are pharmacokinetically important properties when searching for a new drug. The BOILED-Egg graph was used to predict the penetration by computing the lipophilicity and polarity of the molecules. In the BOILED-Egg diagram (Figure 6), this complex clearly differed from the other coordination compounds. Servis ProTox II classified compounds $C_1$, $C_2$, and $C_4$ into toxicity class 4 (predicted LD50: 1000 mg/kg for $C_1$; predicted LD50: 750 mg/kg for $C_2$; predicted LD50: 450 mg/kg for $C_4$) and compound $C_3$ into toxicity class 5 (predicted LD50: 2257 mg/kg).
4. Conclusions

Four complexes of Cd(II) with benzimidazole derivatives were successfully synthesized and named C1, C2, C3, and C4. The composition of the obtained complexes was confirmed by using FAAS, FTIR, and MS coupled with a TG study. All compounds were hydrates. After dehydration, non-conjugated pyridine ring degradation took place. The next step of ligand decomposition was accompanied by the partial removal of the cadmium
content in the form of cadmium chloride, a random amount of which evaporated above its melting point. All of the results obtained from the thermogravimetric analysis and mass spectrometry were compatible with the respective structures of the obtained compounds and their properties. It can be concluded that, in the case of the $C_1$ and $C_3$ compounds, the coordination took place through the nitrogen atom contained in the pyridine ring [47]. Compounds $C_2$ and $C_4$, which had high activity, had a very similar coordination. Cadmium coordinates through the nitrogen atom of the five-membered ring. The results obtained confirmed that the conversion of the ligands into the respective metal complexes significantly improved their anticancer properties. All of them showed much stronger activity than that of etoposide, which is a component of therapy for glioblastoma, neuroblastoma, and lung cancer. The most promising Cd (II) complex, $C_2$, inhibited the growth of neuroblastoma cells five times more than etoposide, with an $IC_{50}$ equal to 22.46 µM (vs. 115.25 µM for etoposide). The antibacterial efficiency of the complexes against Gram-positive bacteria decreased in the following order: $C_1 > C_2 > C_4 > C_3$. Notably, all tested Cd complexes revealed good activity against yeasts in comparison to their ligands. The obtained compounds showed very good antitumor properties, which will allow for future research on the mechanisms of their binding to a particular target or active site (enzyme, protein), as well as their potential use as cytostatic drugs.

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References
1. Kauffman, S. Homeostasis and Differentiation in Random Genetic Control Networks. *Nature* **1969**, *224*, 177–178. [CrossRef] [PubMed]
2. Hanna, N.; Einhorn, L.H. Testicular cancer: A reflection on 50 years of discovery. *J. Clin. Oncol.* **2014**, *32*, 3085–3093. [CrossRef] [PubMed]
3. Dasari, S.; Tchounwou, P.B. Cisplatin in cancer therapy: Molecular mechanisms of action. *Eur. J. Pharmacol.* **2014**, *740*, 364–378. [CrossRef] [PubMed]
4. Cortez, A.J.; Tudrej, P.; Kujawa, K.A.; Lisowska, K.M. Advances in ovarian cancer therapy. *Cancer Chemother. Pharmacol.* **2018**, *81*, 17–38. [CrossRef] [PubMed]
5. Jayson, G.C.; Kohn, E.C.; Kitchener, H.C.; Ledermann, J.A. Ovarian cancer. *Lancet* **2014**, *384*, 1376–1388. [CrossRef]
6. Farooq, M.A.; Aquib, M.; Farooq, A.; Haleem Khan, D.; Joelle Maviah, M.B.; Sied Filli, M.; Kesse, S.; Boakye-Yiadom, K.O.; Mavlyanova, R.; Parveen, A.; et al. Recent progress in nanotechnology-based novel drug delivery systems in designing of cisplatin for cancer therapy: An overview. *Artif. Cells Nanomed. Biotechnol.* **2019**, *47*, 1674–1692. [CrossRef]
7. Vermorken, J.B.; Mesia, R.; Rivera, F.; Remenar, E.; Kawecki, A.; Rottey, S.; Erfan, J.; Zabolotnyy, D.; Kienzer, H.-R.; Cupissol, D.; et al. Platinum-Based Chemotherapy plus Cetuximab in Head and Neck Cancer. *N. Engl. J. Med.* **2008**, *359*, 1116–1127. [CrossRef]
8. Carbone, D.P.; Reck, M.; Paz-Ares, L.; Creeblan, B.; Horn, L.; Steins, M.; Felip, E.; van den Heuvel, M.M.; Cuileanu, T.-E.; Badin, F.; et al. First-Line Nivolumab in Stage IV or Recurrent Non–Small-Cell Lung Cancer. *N. Engl. J. Med.* **2017**, *376*, 2415–2426. [CrossRef]
9. Mok, T.S.K.; Wu, Y.L.; Kudabba, I.; Kowalski, D.M.; Cho, B.C.; Turna, H.Z.; Castro, G.; Srimuninnimit, V.; Laktionov, K.K.; Bondarenko, I.; et al. Pembrolizumab versus chemotherapy for previously untreated, PD-L1-expressing, locally advanced or metastatic non-small-cell lung cancer (KEYNOTE-042): A randomised, open-label, controlled, phase 3 trial. *Lancet* **2019**, *393*, 1819–1830. [CrossRef]
10. Farrell, N. Metal Complexes as Drugs and Chemotherapeutic Agents. *Compr. Coord. Chem. II* **2004**, *9*, 809–840. [CrossRef]
11. Johnson, N.P.; Butour, J.-L.; Villani, G.; Wimmer, F.L.; Defais, M.; Pierson, V.; Brabc, V. Metal Antitumor Compounds: The Mechanism of Action of Platinum Complexes. Prog. Clin. Biochem. Med. 1989, 10, 1–24. [CrossRef]

12. Ceremosimo, R.J. Cisplatin neurotoxicity. Cancer Treat. Rev. 1989, 16, 195–211. [CrossRef]

13. Leo, M.; Schmitt, L.I.; Erkel, M.; Melnikova, M.; Thomale, J.; Hagenacker, T. Cisplatin-induced neuropathic pain is mediated by upregulation of N-type voltage-gated calcium channels in dorsal root ganglion neurons. Exp. Neurol. 2017, 288, 62–74. [CrossRef]

14. Chen, S.H.; Chang, J.Y. New insights into mechanisms of cisplatin resistance: From tumor cell to microenvironment. Int. J. Mol. Sci. 2019, 20, 4136. [CrossRef]

15. Galluzzi, L.; Vitale, I.; Michels, J.; Brenner, C.; Szabadkai, G.; Harel-Bellan, A.; Castedo, M.; Kroemer, G. Systems biology of cisplatin resistance: Past, present and future. Cell Death Dis. 2014, 5, e1257. [CrossRef]

16. Rocha, C.R.R.; Silva, M.M.; Quinet, A.; Cabral-Neto, J.B.; Menck, C.F.M. DNA repair pathways and cisplatin resistance: An intimate relationship. Clinics 2018, 73, e478s. [CrossRef]

17. Alloway, B.J. Heavy Metals and Metalloids as Micronutrients for Plants and Animals. In Heavy Metals in Soils, Environmental Pollution; Springer Science & Business Media: Berlin/Heidelberg, Germany, 2013; pp. 195–209. [CrossRef]

18. IARC. International Agency for Research on Cancer Evaluation of Carcinogenic. Glass 1993, 58, 119–130.

19. Agency for Toxic Substances and Disease Registry Division of Toxicology and Human Health Sciences. Support Document to the 2019 Substance Priority List (Candidates for Toxicological Profiles); Agency for Toxic Substances and Disease Registry Division of Toxicology and Human Health Sciences of the U.S. Department of Health and Human Services: Washington, DC, USA, 2019; pp. 1–12.

20. Panchsheela Ashok, U.; Prasad Kollur, S.; Prakash Arun, B.; Sanjay, C.; Shrikrishna Suresh, K.; Anil, N.; Vasant Baburao, H.; Markad, D.; Ortega Castro, J.; Frau, J.; et al. In vitro anticancer activity of 4(3H)-quinazolinolone derived Schiff base and its Cu(II), Zn(II) and Cd(II) complexes: Preparation, x-ray structural, spectral characterization and theoretical investigations. Int. J. Inorg. Chem. 2020, 511, 119846. [CrossRef]

21. Ye, C.H.; Chen, G.; Gong, Y.L. Two Heteroligand Cd(II)-coordination Polymers: Crystal Structures and Anti-Lung Cancer Activity Evaluation. Russ. J. Coord. Chem. Khimiya 2020, 46, 653–661. [CrossRef]

22. Icsel, C.; Yilmaz, V.T.; Aydinlik, S.; Aygun, M. Zn(ii) and Hg(ii) saccharinate complexes with 2,6-bis(2-benzimidazolyl)pyridine and ethylenediamine. Chem.-An Asian J. 2019, 14, 4420–4428. [CrossRef]

23. Icsel, C.; Yilmaz, V.T.; Aydinlik, S.; Aygun, M. Zn(ii) and Hg(ii) saccharinate complexes with 2,6-bis(2-benzimidazolyl)pyridine and ethylenediamine. Chem.-An Asian J. 2019, 14, 4420–4428. [CrossRef]

24. Chen, Y.Q.; Tian, Y.; Yao, S.L.; Zhang, J.; Feng, R.Y.; Bian, Y.J.; Liu, S.J. CdII-Organic Frameworks Fabricated with a N-Rich Ligand and Flexible Dicarboxylates: Structural Diversity and Multi-Responsive Luminescent Sensing for Toxic Anions and Ethylenediamine. Int. J. Mol. Sci. 2016, 17, 2014. [CrossRef]

25. Markad, D.; Ortega Castro, J.; Frau, J.; et al. In vitro anticancer activity of 4(3H)-quinazolinolone derived Schiff base and its Cu(II), Zn(II) and Cd(II) complexes: Preparation, x-ray structural, spectral characterization and theoretical investigations. Int. J. Inorg. Chem. 2020, 511, 119846. [CrossRef]

26. Krasnovskaya, O.; Naumov, A.; Guk, D.; Gorelkin, P.; Erofeev, A.; Beloglazkina, E.; Majouga, A. Copper coordination compounds as biologically active agents. Int. J. Mol. Sci. 2020, 21, 3965. [CrossRef]

27. Ye, C.H.; Chen, G.; Gong, Y.L. Two Heteroligand Cd(II)-coordination Polymers: Crystal Structures and Anti-Lung Cancer Activity Evaluation. Russ. J. Coord. Chem. Khimiya 2020, 46, 653–661. [CrossRef]

28. Connell, T.U.; Schieber, C.; Silvestri, I.P.; White, J.M.; Williams, S.J.; Donnelly, P.S. Copper coordination compounds as biologically active agents. Int. J. Mol. Sci. 2020, 21, 3965. [CrossRef]

29. Plumb, J.A. Cell sensitivity assays: Clonogenic assay. Methods Mol. Med. 2004, 88, 159–164. [CrossRef]

30. Plumb, J.A. Cell sensitivity assays: Clonogenic assay. Methods Mol. Med. 2004, 88, 159–164. [CrossRef]

31. Jin, X.; Chen, H.; Zhang, W.; Wang, B.; Shen, W.; Lu, H. A novel purine derivative-based colorimetric chemosensor for sequential detection of copper ion and sulfide anion. J. Am. Chem. Soc. 2020, 142, 2805–2814. [CrossRef]

32. Girek, M.; Kłosiński, K.; Grobelski, B.; Pizzimenti, S.; Cucci, M.A.; Daga, M.; Barrera, G.; Pasieka, Z.; Czarnecka, K.; Szymański, P. Novel tetrahydroacridine derivatives with iodobenzoic moieties induce G0/G1 cell cycle arrest and apoptosis in A549 non-small lung cancer and HT-29 colorectal cancer cells. Mol. Cell. Biochem. 2019, 460, 123–150. [CrossRef]

33. Daina, A.; Michielin, O.; Zoete, V. SwissADME: A free web tool to evaluate pharmacokinetics, drug-likeness and medicinal chemistry friendliness of small molecules. Sci. Rep. 2017, 7, 24717. [CrossRef]

34. Daina, A.; Michielin, O.; Zoete, V. iLOGP: A Simple, Robust, and Efficient Description of n-Octanol/ Water Partition Coefficient for Drug Design Using the GB/SA Approach. J. Chem. Inf. Model. 2014, 54, 3284–3301. [CrossRef] [PubMed]

35. Daina, A.; Zoete, V. A BOILED-Egg to Predict Gastrointestinal Absorption and Brain Penetration of Small Molecules. ChemMedChem 2016, 11, 1117–1121. [CrossRef] [PubMed]

36. Banerjee, P.; Eckert, A.O.; Schrey, A.K.; Preisssner, R. ProTox-II: A webserver for the prediction of toxicity of chemicals. Nucleic Acids Res. 2018, 46, W257–W263. [CrossRef] [PubMed]

37. Raducka, A.; Czyłkowski, A.; Gobis, K.; Czarnecka, K.; Szymański, P.; Świątkowski, M. Characterization of metal-bound benzimidazole derivatives, effects on tumor cells of lung cancer. Materials 2021, 14, 2958. [CrossRef] [PubMed]
37. Kruszynski, R.; Sierań ski, T.; Święatkowski, M.; Zielak, M.; Wojciechowski, J.; Dzierzawska, M.; Lewiński, B. On the coordination behavior of the hmta toward zinc and cadmium cations in presence of sulfate(VI) and nitrate(V) anions. J. Coord. Chem. 2014, 67, 1332–1352. [CrossRef]
38. Gu, J.; Wan, S.; Kirillova, M.; Kirillov, A. H-bonded and metal(II)-organic architectures assembled from an unexplored aromatic tricarboxylic acid: Structural variety and functional properties. Dalt. Trans. 2020, 49, 7197–7209. [CrossRef] [PubMed]
39. Rosenstock, H.M.; Stockbauer, R.; Parr, C. Unimolecular kinetics of pyridine ion fragmentation. Int. J. Mass Spectrom. Ion Phys. 1981, 38, 323–331. [CrossRef]
40. Lide, D.R. Standard density of water. In CRC Handbook of Chemistry and Physics, Internet Version 2007, 87th ed.; Taylor and Francis: Boca Raton, FL, USA, 2007; pp. 1–2317.
41. Kruszynski, R. Synthesis of coordination compounds via dehalogenation of zinc bromoacetate in presence of some amines. Int. J. Inorg. Chem. 2011, 371, 111–123. [CrossRef]
42. Lipinski, C.A.; Lombardo, F.; Dominy, B.W.; Feeney, P.J. Experimental and computational approaches to estimate solubility and permeability in drug discovery and development settings. Adv. Drug Deliv. Rev. 2012, 64, 4–17. [CrossRef]
43. Ghose, A.K.; Viswanadhan, V.N.; Wendoloski, J.J. A knowledge-based approach in designing combinatorial or medicinal chemistry libraries for drug discovery. 1. A qualitative and quantitative characterization of known drug databases. J. Comb. Chem. 1999, 1, 55–68. [CrossRef]
44. Egan, W.J.; Merz, K.M.; Baldwin, J.J. Prediction of drug absorption using multivariate statistics. J. Med. Chem. 2000, 43, 3867–3877. [CrossRef]
45. Veber, D.F.; Johnson, S.R.; Cheng, H.Y.; Smith, B.R.; Ward, K.W.; Kopple, K.D. Molecular properties that influence the oral bioavailability of drug candidates. J. Med. Chem. 2002, 45, 2615–2623. [CrossRef]
46. Muegge, I.; Heald, S.L.; Brittelli, D. Simple selection criteria for drug-like chemical matter. J. Med. Chem. 2001, 44, 1841–1846. [CrossRef]
47. Raducka, A.; Święatkowski, M.; Korona-Głowniak, I.; Kaproń, B.; Plech, T.; Szczesio, M.; Gobis, K.; Szynkowska-Jóźwik, M.I.; Czylkowska, A. Zinc Coordination Compounds with Benzimidazole Derivatives: Synthesis, Structure, Antimicrobial Activity and Potential Anticancer Application. Int. J. Mol. Sci. 2022, 23, 6595. [CrossRef]