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Cellular uptake of metallated cobalamins

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Cellular uptake of vitamin B12-cisplatin conjugates was estimated via detection of their metal constituents (Co, Pt, and Re) by inductively coupled plasma mass spectrometry (ICP-MS). Vitamin B12 (cyano-cob(III)alamin) and aquo-cob(II)alamin [Cbl-OH2]+, which differ in the β-axial ligands (CN− and H2O, respectively), were included as control samples. The results indicated that B12 derivatives delivered cisplatin to both cellular cytosol and nuclei with an efficiency of one third compared to the uptake of free cisplatin cis-[PtCl2(NH3)2]. In addition, uptake of charged B12 derivatives including [Cbl-OH2]+, [(Co-CN-(cis-PCl(NH3)2)]+, [(Re)-(Co-CN-(cis-PCl(NH3)2)]+, and [(Co-CN-(trans-Pt(Cyt)(NH3)2)]2+ (Cyt = cytarabin) was high compared to neutral B12, which implied the existence of an additional internalization pathway for charged B12 vitamin analogs. The affinities of the charged B12 derivatives to the B12 transporters HC, IF and TC were similar to that of native vitamin B12.

Introduction

Cyano-cob(III)alamin, also designated vitamin B12, is an essential, water-soluble, and non-toxic vitamin even at high doses. Vitamin B12 is involved in the methionine synthesis and the tricarboxylic acid cycle. B12 comprises an octahedral CoIII complex with a tetradentate corrin ring, an α-nucleotide loop and a CN− ligand in the β-axial position. The cyanide ligand can be exchanged for other monodentate functions (Fig. 1).1 Intracellular reduction of the CoIII center subsequently forms square pyramidal cob(III)alamin and square planar cob(II)alamin, respectively.2,3 Enzymatic alkylation of cob(II)alamin in the cytosol and mitochondria leads to the formation of two B12-coenzymes with methyl and adenosyl as β-ligands.4,5 B12 is protected from degradation inside living systems through its binding to the proteins haptocorrin (HC, present in saliva/gastric fluids), intrinsic factor (IF, secreted by parietal cells), and transcobalamin (TC, plasma protein).6,7 Binding of B12 to the proteins haptocorrin (HC, present in saliva/gastric fluids), intrinsic factor (IF, secreted by parietal cells), and transcobalamin (TC, plasma protein).8–11 Numerous functional groups on the corrin side chains, β-ligand, and α-nucleotide loop (Fig. 1) are available to coordinate or conjugate drugs and analytical modalities.12 The β-axial ligand is a preferred derivatization site owing to its ease-of-release in cellular environment and its little involvement in interaction with proteins. Nitrogen mustard chlorambucil,13 colchicine,14 cisplatin,15 and insulinomimetic vanadate (V5) compound16 are drugs which have been bound to B12 at this site. In addition, fluorophores17 and radionuclides18,19 have also been attached to B12 at this position. The ribose-hydroxyl group is another convenient derivatization site as it is not involved in protein interaction.20 Fluorophores,21,22 cytotoxic drugs,23 insulin,24 and appetite suppressing peptide hPYY25 have also been coupled to B12 via this position.

Derivatization of B12, however, is not limited to the β- and ribose-positions. Radioactive 111In-DTPA and 99mTc-DTPA were coupled to B12 via the b-side chain.26,27 The 99mTc(CO)3 moiety was coupled to B12 via ligands coupled to the b- and d-side chains28 and biotin via either the b-, c-, d-, or e-side
chains. Increasing structural knowledge of B12 and of its transport proteins is the base for fine-tuning the structures of biologically active B12 derivatives and mimetics. Several B12 antagonists were reported, including B12-ε-lactam, and B12 with de novo peptide backbones. In addition, tissue distribution was interfered by modifying the B12 structure at a critical site (b-side chain) for protein interaction.

Research programs have been dedicated to explore the correlation of B12 requirements and DNA synthesis. This, however, is not the only advantage of using B12 as a carrier for anti-cancer drugs. Its water solubility allows convenient administration of the attached insoluble drugs. B12 protects for anti-cancer drugs. Its water solubility allows convenient administration of the attached insoluble drugs. B12 protects against the regime.

Results and discussion

Synthesis of $([\text{Re}-\text{Co}-\text{CN}-(\text{cis-PtCl(NH}_3)_2])$)

The tridentate NNO ligand (6) was chelated to the $\{\text{fac-Re(CO)}_3\}^+$ moiety prior to conjugation to the B12 structure (step iv and v), see Scheme 1. In the next step, the $\beta$-CN $^-$ was coordinated to $[\text{cis-PtCl(OH}_2)(\text{NH}_3)_2]^+$ by substituting the aquo-ligand bound to PtII (step vi) to yield 2. This synthesis scheme produced a high yield (80%) of $[\text{Re}-\text{Co}-\text{CN}-(\text{cis-PtCl(NH}_3)_2])$ (2, Scheme 1).

The IR stretching frequency of CN $^-$ was higher in 2 (2200 cm$^{-1}$) than in B12 (2134 cm$^{-1}$). It confirmed the formation of $\{\text{Co}}–\text{CN–{Pt} bonds.}$ Compared to the 1H NMR spectrum of $[\text{Co}-\text{CN}-(\text{cis-PtCl(NH}_3)_2])$ previously published shows four additional 1H signals (from the Re-[chelate]-ligand) in the fingerprint aromatic region ($>$ 5.5 ppm (unmarked peaks in Fig. 3), which further confirmed the presence of the rhenium complex.

Affinity of B12 derivatives towards transport proteins

Affinity to transport proteins of B12 derivatives was measured by Differential Scanning Fluorometry (DSF), where a fluorescent dye interacts with the hydrophobic sites of unfolded proteins and the melting temperature ($T_m$) of proteins can be inferred from the recorded fluorescence intensity. Following ligand
binding, the protein stability is enhanced and the ligand-bound proteins unfold at higher temperature with respect to the free proteins. B12 derivatives which bind to B12 transport proteins would produce comparable \( \Delta T_m \) values to those of B12 while \( \Delta T_m \) values would be much lower if these derivatives do not bind to B12 transport proteins. In Fig. 4, B12 was used as a positive control for the three transport proteins and cobinamide (Cbi) was used as a negative control. Despite the chemical modification of the B12 structure, compound 2 retained good affinity to all B12 transport proteins as it has comparable \( \Delta T_m \) values to those of B12.

Cellular uptake of \([\text{Co-CN-}(\text{cis-PtCl(NH}_3)_2])^+\), \([\text{Re}-\text{Co}-\text{CN-}(\text{cis-PtCl(NH}_3)_2])^+\) and \([\text{Co-CN-}(\text{trans-Pt(Cyt)(NH}_3)_2])^2+\)

To obtain information on the action of these compounds, knowledge on the amount of drug taken up by cells and accumulated in the nucleus with and without B12 is required and is the base for evaluating the efficiency of B12 as a tumor-targeting drug carrier molecule. The metals (Co, Re, Pt) in B12-drugs conjugates can be measured by ICP-MS, i.e., cytosolic and nucleic determination of \(^{50}\text{Co},\ ^{195}\text{Pt},\) and \(^{187}\text{Re}\) reveal the intracellular distribution of these compounds. Fig. 5A shows the cellular cisplatin uptake in K562 cells, determined as the content in the cytosol. The uptake of platinum containing B12 derivatives (1, 2, and 3) is approximately one third compared to the uptake of cisplatin. Similarly it is seen from Fig. 5B that the platinum uptake into cellular nuclei of 1, 2, and 3 was low compared to cisplatin. The low nucleic platinum uptake of 3 was expected as it has a transplatin-based structure. More interestingly, substantial extracellular Pt-release from the B12 structures, which would have resulted in comparative cellular uptake to cisplatin, was not observed. These data indicate that Pt is not released in the extracellular media but taken up together with B12. The complexes are protected when conjugated to the B12 structure. Unexpectedly, the cytosolic uptake of B12 was considerably lower than of \([\text{Cbl-OH}_2]^+\), 1, 2, and 3 (Fig. 5). This indicated that the observed uptake was not mediated by TC as this protein has similar affinities to B12, \(2\) (Fig. 4), and \(3\). Therefore TC would have allowed similar accumulation of these compounds in cellular components.

It was previously reported that B12 derivatives are able to enter cells without the support of TC. This uptake was, nevertheless, less efficient. \(^{45-48}\) HeLa cells internalized only 1–2% of free B12 with respect to holo-TC. \(^{45}\) It was later concluded that free B12 supported cell growth ca. 100 to 1000-fold less efficiently than holo-TC. \(^{49}\) However, human skin fibroblasts could internalize free B12 better than holo-TC, up to more than 20% were reported. \(^{46}\) Berliner et al. proposed a free B12-specific receptor on the cell membrane \(^{46}\) while Hitzig et al. suggested an α2-globulin-like protein in serum as the alternative pathway for cell to internalize free B12 derivatives. \(^{50}\)

In this study, we observed higher accumulation of \([\text{Cbl-OH}_2]^+\) as compared to B12. In a previous report, it was found that \([\text{Cbl-OH}_2]^+\) was internalized and retained better than B12 in human skin
fibroblasts culture but the root of this difference was not fully understood. As we also found comparable cellular uptake among compounds 1, 2, 3, and [Cbl-OH2], we conclude that the positive charge carried by these compounds allows cells to internalize them better than the neutral B12 derivatives (Fig. 5). Behavioral differences between charged and neutral B12 derivatives were also previously observed. It is possible that charged B12 compounds were either distinctly recognized by a B12-specific receptor or they simply passed through the cationic channels on cell membranes. This, however, requires further clarification. In any case, the experimental results highlighted once more the complex multichannel uptake pathways of B12 vitamin analogues including internalization of holo-TC, free B12 derivatives, and charged B12 derivatives.

The preferable cellular import of charged B12 compounds should be kept in perspective for future design of pharmaceutical B12 derivatives.

Drugs like cisplatin and cytarabine attached to the β-CN− of B12 are released by reducing enzymes in the cytosol. Furthermore, subsequent cytotoxic activities of the released drugs from [(Co-CN-(cis-PtCl(NH3)2))]+ (1) and [(Co-CN-(trans-Pt(Cyt)(NH3)2)]z+ (3) were also demonstrated. Cisplatin interferes with cell division by binding to DNA. Previous studies demonstrated accumulation of cisplatin in the cytosol and the nucleus in non-adherent EATC and adherent ELA cells. It is therefore especially important for compound 1 and 2 to deliver platinum to nucleus for DNA interaction. As evident from Fig. 6, platinum accumulated in the nucleus in EATC and in ELA cells similar to K562 cells. Furthermore, we found that the absolute amount of platinum bound to DNA in cells exposed for 18 hours to 10 μM and 50 μM [(Re)-(Co-CN-(cis-PtCl(NH3)2)]− (2) was 19 and 83 pmol μg−1 DNA (EATC) and 5 and 20 pmol μg−1 DNA (ELA). Hence, cisplatin coupled to B12 derivative did accumulate in the nucleus and was bound to DNA.

The uptake of Pt into cellular components was always higher than Co (Fig. 5 and 6). A correlation between cobalt and platinum uptake for the three compounds 1, 2 and 3 is shown in Fig. 7. The equation for the trend line is y = 1.60X + 4.9. The one outlier point for EATC, cytosol was omitted. The correlation shows that platinum is taken up by a factor of 1.6 higher than cobalt as estimated from the cytosolic and nucleic content.

Conclusions

We reported here comparable uptake in cellular nuclei and cytosol of different B12–drug conjugates, including [(Co-CN-(cis-PtCl(NH3)2)]+, [(Re)-(Co-CN-(cis-PtCl(NH3)2)]−, and [(Co-CN-(trans-Pt(Cyt)-(NH3)2)]z+. The results underline that these B12 derivatives deliver cisplatin and cisplatin-related drugs to both, cytosol and nuclei, in two different cancer cell lines. It was found that charged B12 derivatives were preferentially taken up into cellular compartments as compared to neutral and native B12. We therefore propose the existence of an additional uptake pathway for charged B12 derivatives in addition to the previously known TC-mediated B12 uptake pathway. Accumulation of released platinum in the nucleus was clearly observed and explains the cytotoxic action of these compounds as previously described.

Experimental

Materials and methods

Analytical high performance liquid chromatography (HPLC) analyses were performed using a Merck-Hitachi L-7000 system...
equipped with a diode-array UV/Vis spectrometer. Reaction progresses to 2, 3, 7 and 8 were monitored by HPLC. Preparative HPLC purification was performed on Varian Prostar system with two Prostar 215 pumps and a Prostar 320 UV/Vis detector. Reaction progresses to 4, 5, and 6 were assessed by TLC. After purification the B12 derivatives were lyophilized. Purity was assessed by analytical HPLC, IR, ESI-MS (Bruker Daltonic) and NMR. NMR spectra were recorded by Bruker DRX 500 MHz spectrometer. Chemical shifts are reported relative to residual solvent protons and carbons. The chemical shifts of 195Pt NMR spectra were relative to Na2PtCl6 in D2O.

Phosphate-buffered saline (PBS) contained 137 mM NaCl, 2.6 mM KCl, 6.5 mM NaH2PO4, and 1.5 mM KH2PO4. Lysis buffer was prepared from 150 mM NaCl, 20 mM HEPES buffer, 1 mM EDTA, 0.5% Triton X-100 (Pharmacia Biotech AB), 1% protease inhibitor cocktail (freshly added), 1 mM NaN3 (freshly added). All aqueous solutions for ICPMS measurement were prepared in purified water (>18.2 MΩ) from Milli-Q deionization unit. Standards and samples were prepared in a mixture of HNO3 0.65% : HCl 0.1%.

HPLC conditions. The analytical HPLC column was Macherey Nagel Nucleosil C18 (5 μm, 100 Å, 250 mm × 3 mm). The preparative HPLC column was Macherey Nagel Nucleosil C18 (7 μm, 100 Å, 250 mm × 40 mm) with a flow rate of 35 ml per minute. The following solvent system was used: 0.1% trifluoroacetic acid (TFA) (solvent A), methanol (MeOH) (solvent B). The following solvent gradient was used: gradient 1, 0–5 minutes: 0% A, 75% A, 5–30 minutes: 0% A.

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Syntheses

N-tert-Butyloxycarbonyldiamines (4), 34 [NEt3][ReBr6(CO)3] 33 and [[Co-CN-trans-Pr(Cy)(NH)2]3] 34 were prepared according to literature procedures. Compounds (5) and (6) were prepared as described in literature 35 but with the following modifications: 2 g (4) (11 mmol) was dissolved in 100 ml MeOH. 1.3 g pyridine-2-carbaldehyde (12 mmol) was added dropwise. After stirring for 3 hours at room temperature, the solvent was removed and the imine dissolved in 50 ml ethanol (EtOH). Reduction with 1.5 g NaNBH4 in 2 h, removal of solvent, re-dissolution in diluted NaHCO3 and extraction into CH2Cl2 gave the raw product. The purification of 5 was performed with silica gel column (ethylacate, methanol 1:3) (2.8 g, 91%). 1H-NMR (200 MHz, MeOD, δ in ppm): δ = 8.63 (m, 1H), 7.87 (m, 1H), 7.49 (m, 1H), 7.41 (m, 1H), 6.58 (s, 1H), 6.26 (s, 1H), 5.33 (s, 1H), 4.57 (d, J = 16 Hz, 1H), 4.40 (d, J = 15.5 Hz, 1H), 3.70 (m, 2H), 3.08 (m, 2H), 1.81 (m, 2H), 1.48 (m, 2H), 1.4 (s, 1H). IR (KBr, cm⁻¹): 3436, 3393, 3283, 2920, 1762, 1566, 1370, 1161, 1152, 1068, 1000, 824. ESI-MS: [M + H]⁺ = 364.3 [M + Na⁺].

Synthesis of (8). 50 mg (82.5 μmol) of (7) was Boc deprotected with 1 ml TFA in 3 ml CH2Cl2. To a solution of 150 mg (0.11 mmol) B12 in 3.5 ml dry DMSO was added 24 mg (0.15 mmol) CDT. The mixture was stirred for 1 h at 60 °C. Then the Boc-deprotected product was added together with 4 ml dry DMSO and 0.8 ml triethylamine. The reaction was stirred overnight. Diethyl ether was added to precipitate the product. Separation of (8) was performed on preparative HPLC, solvent system 1, gradient 1 (31 mg, 14.5%). 1H-NMR (500 MHz, MeOD, δ in ppm): δ = 8.1 (d, J = 5.5 Hz, 1H), 8.07 (m, 1H), 7.71 (t, J = 6.5 Hz, 1H), 7.52 (m, 1H), 7.25 (s, 1H), 7.13 (d, J = 4 Hz, 1H), 6.57 (s, 1H), 6.24 (s, 1H), 5.97 (s, 1H). IR (KBr, cm⁻¹): 2136 (CN), 2029, 1918, 1902 (CO). ESI-MS: [M + H]⁺ = 1888 [M + H⁺], 945 [M + 2H⁺]²⁺.

Synthesis of (2). 12 mg (40 μmol) of cisplatin and 6.7 mg (39 μmol) of AgNO3 were mixed in 1 ml H2O. The reaction was stirred in the absence of light at 40 °C for 4 h. AgCl was filtered and the filtrate was added to 22 mg (11.6 μmol) of (8). The reaction was stirred at 45 °C for one day in the absence of light. The product was purified by preparative HPLC, solvent system 1, gradient 1 (20 mg, 80%). 1H-NMR (500 MHz, MeOD, δ in ppm): δ = 8.8 (d, J = 5 Hz, 1H), 8.08 (d, J = 6 Hz, 1H), 7.72 (m, 1H), 7.54 (m, 1H), 7.25 (s, 1H), 7.14 (s, 1H), 6.58 (s, 1H), 6.26 (s, 1H), 5.99 (s, 1H). 195Pt-NMR (107 MHz, MeOD, δ in ppm): δ = -2354. IR (KBr, cm⁻¹): 2200 (CN), 2026, 1916, 1904 (CO). ESI-MS: [M + H]⁺ = 1076.6 [M + 2H⁺]²⁺.

Affinity to transport proteins

Binding to HC/IF and TC was measured by differential scanning fluorimetry (DSF). Details of the method, protein expression and purification was previously described. 42

Uptake measurement with ICPMS. Cells were seeded in T-75 flasks with a seeding density of 3 × 10⁵ cells per flask. The culture medium was 12 ml Roswell Park Memorial Institute (RPMI)-1640 with 10% fetal bovine serum (FCS) and penicillin-streptomycin (100 U ml⁻¹). After 1 day of incubation, each flask was treated with 10 μM of a compound. These included cisplatin, B123, [Cbl-OH]⁺, (1), (2), and (3). Cells were further maintained in culture for two days. Nucleic and cytosolic...
fractions were separated according to a previously published procedure with certain modifications. In details, cells were centrifuged for 1 minute at 700 g and room temperature (adherent ELA cells were trypsinized with 0.25% trypsin/EDTA in PBS before this step). The supernatant was discarded. Pellets were washed (3 x) with 5 ml PBS before chemical lysis. Lysis buffer was added. 20 μl of the lysis was spared for protein determination (DC Protein Assay, Bio-Rad). Remaining lysis was centrifuged at 600 g, 4 °C for 5 minutes. The pellet was washed (3 x) with 1 ml NaCl/KCl (30 mM/120 mM) solution to isolate nuclei fraction. The supernatant was centrifuged at 5500 g, 4 °C for 15 minutes. Cytosolic fraction was the supernatant part. The nuclei fractions were digested with 100 μl concentrated HNO3.

DNA was isolated according to the following procedure. Cells were centrifuged two times for 5 s at 13 000–16 000 g. The precipitate was subsequently lysed with 300 μl lysis solution. 100 μl of Protein Precipitation Solution was added and mixed well. After the proteins were precipitated, the supernatant was discarded. Pellets were washed (3 x) with 1 ml DNA hydration solution. 100 μl concentrated HNO3 (50 μl) was then added to the dry DNA. The mixture was then incubated in a water bath at 65 °C for 1 hour to dissolve DNA, followed by incubating at room temperature overnight. Both subcellular fractions were diluted with 0.65% HNO3/0.1% HCl for ICPMS measurement.

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