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6-O-Nucleotidyltransferase: an aminoglycoside-modifying enzyme specific for streptomycin/streptidine

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antibiotics

Aminoglycosides are especially useful for the treatment of hospital-acquired infections. The main problem for the application of these antibiotics is the presence of bacterial resistance enzymes, in particular, nucleotidyltransferases (ANTs). These enzymes catalyze the transfer of an adenylyl group from the MgATP complex to different positions of antibiotic. To understand the mechanisms that lead to antibiotic inactivation, we have performed a comprehensive experimental analysis of one of those enzymes. The 6-O- nucleotidyltransferase enzyme (ANT(6)) from Bacillus subtilis voc cloned, overexpressed and purified in E. coli. The kinetic parameters revealed a narrow specificity of the ANT(6) for MgATF /streptomycin as substrates. The binding epitope of the streptomycin recognized by the ANT(6) is the streptidine moiety Therefore, the use of streptidine as a "decoy acceptor" allows the recovery of the antibiotic activity of streptomycin E. coli cells that are overexpressing the ANT(6).

resistance

Introduction

Aminoglycoside antibiotics are of special interest in medicine for the treatment of severe bacterial infections.¹⁻³ These antibiotics are active against either Gram+ or Gram- bacteria, since their primary mechanism of action is the inhibition of bacterial protein synthesis by binding to the tRNA acceptor A site in the small ribosomal subunit that is causing a misreading and/or hindering the translocation step.⁴⁻⁸ For most of the aminoglycosides, the aminocyclitol moiety is а 2deoxystreptamine ring (2-DOS) and can be substituted at positions 4 and 6 (kanamycin family (3)) or at 4 and 5 (neomycin (2) family). However, a number of active aminoglycosides are structurally atypical, such as streptomycin (ST) (1) (Fig. 1).

Keywords:

aminoglycosides

The emergence of bacterial resistance for all classes of antipathogenic agents has become a serious problem over recent years.⁹⁻¹¹ Aminoglycosides were one of the first groups of antibiotics to meet the challenge of resistance.¹² The most prevalent source of clinically relevant resistance to aminoglycoside antibiotics is conferred by the enzymatic inactivation of the drugs by the aminoglycoside-modifying enzymes (AMEs).^{13,14}

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The AMEs can be classified as N-acetyltransferases (AACs), Oadenyl transferases (ANTs) or O-phosphotransferases (APHs) In each of these families, there are several enzymes that catalyzed the reactions with different regioselectivities and substrate specificities.

enzymes

decov



Fig. 1. Structures of aminoglycoside antibiotics: streptomycin (1), neomy (2) and kanamycin A (3).

The ANT' family is the smallest of the three groups with only a few enzymes identified to date [ANT(6)], [ANT(9)], [ANT(4')]

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⁺ Footnotes relating to the title and/or authors should appear here. Electronic Supplementary Information (ESI) available: [details of any supplementary information available should be included here]. See DOI: 10.1039/x0xx00000x

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[ANT(2")].¹⁵⁻¹⁸ One of the most prevalent ANT enzymes is the streptomycin adenylyltransferase (ANT (6)).¹⁹⁻²¹ This enzyme catalyzes the transfer of an adenyl group from the [MgATP]²⁻ complex to the OH at the 6 position of streptidine unit in the streptomycin (1), thus leading to a sharp decrease in the drug's affinity for its RNA target (Scheme 1). Streptomycin (1) was the first aminoglycoside antibiotic to be discovered in 1943. It has a three-ring structure (Scheme 1), comprising a substituted aminocyclitol (streptidine moiety), linked to a modified ribose (α -L-streptose), which is linked in turn to an N-methyl- α -Lglucosamine residue.



Scheme 1. ANT(6) catalyzes the adenylation of streptidine moiety (red) in streptomycin (1) antibiotic, giving AMP-streptomycin inactive

Some years ago, our group reported that the binding epitope of the streptomycin recognized by this enzyme was the streptidine moiety²²; we also noted its use as a "decoy acceptor" \ddagger23,24 by the ANT(6) enzyme to rescue the antibiotic activity of streptomycin (1) from bacterial strains overexpressing this aminoglycoside-modifying enzyme. We showed that the streptidine was a substrate for the aminoglycoside-inactivating enzyme ANT(6), competing with the streptomycin (1) for the active center of the enzyme. Thus, the addition of this molecule in a cell culture restored the activity of the streptomycin antibiotic normally rendered inactive by the ANT(6) enzyme, because the streptidine was acting as a "decoy acceptor" of the ANT(6).

Therefore, we demonstrated that the use of decoy acceptors can be an effective strategy for combating the phenomenon of resistance promoted by AMEs. However, for the design of more efficient decoy acceptors, a deeper knowledge of the molecular mechanism of the AMEs, their structures and interactions with the drugs is needed. Herein, we report the biochemical characterization of the ANT(6) from Bacillus subtilis. To this end, we cloned, overexpressed the enzyme in E. coli. The kinetic parameters revealed a narrow specificity of the ANT(6) for MgATP /streptomycin as substrates. Finally, we assayed the use of 2-deoxystreptami e as a decoy acceptor, and its ability to restore the antibiotic activity of streptomycin against resistant bacterial strains.

Results and discussion

Cloning, over-expression, purification and characterization of ANT(6)

The *aadK* gene from *B. subtilis* was amplified by PCR, using primers designed specifically to complement 15 bp at the 5'ends of codifying and complementary DNA strains. The recognition sequence for the restriction enzymes EcoRI a HindIII was introduced in the amplification product during the PCR process. The double-digested PCR product was cloned into the vector pET-28b(+) to yield the recombinant plasmid pETaadk NHis₆-tag.

The aadK gene was expressed in E. coli BL21(DE3) at 30 °C. SDS-PAGE analysis of the gene expression showed an IPTG inducible protein of 37.4 kDa of the ANT(6) with an NHis₆-tag After addition of the thrombin enzyme, the ANT(6) AHU enzyme was obtained with 35.5 kDa of weight (Fig. S1). Th€ two-step purification of the ANT(6) yielded well over 40 mg on protein per liter of the cell culture, with a purity degree of a least 95% (Fig. S1). The ANT(6) was aggregated without MgCl₂, so the use of that cation is essential to avoid the formation o aggregations (Fig S2a). The purified ANT(6)∆His is a dimer based on an ultracentrifugation experiment (Fig S2b) and it: mobility over Superdex 200 in standardized analytical-gel filtration experiments

The activity of the pure recombinant enzyme was followed by HPLC and reaction products identified by ¹H-NMR.²⁵ The recombinant ANT(6) exhibited activity between pH 6.5 and pH 8.5, with the maximum activity at pH 7.5. At pH values ovel 7.5, the activity level drops drastically to virtually zero (Fig. S3).

Regarding the secondary structure of the recombinant enzyme, the far-UV CD spectrum of purified ANT(6) shows two minima in ellipticity at ~210 and ~222 nm, indicative of a main α -helical secondary structure (Fig. 2a). The percentages of the different secondary structure elements were estimated using the CDPro²⁶ software package ~48.2% α -helix, ~8.7% β -she t, ~15.9% turns, and ~27.3% of unordered structure we thereby obtained. The tertiary structure of the protein in the microenvironment of the protein fluorophores was analyzed using fluorescence-emission spectroscopy of 19 tyrosines and 7 tryptophan present in the ANT(6) sequence (Fig. 2b). After excitation at 275 nm, the fluorescence spectrum exhibited an emission maximum centered at 345 nm, close to the expected emission of free tryptophans exposed to an aqueous solution (348-350 nm). The emission spectrum obtained up a excitation at 295 nm, was almost identical to the previo spectrum, indicating an almost negligible contribution of tyrosines to the fluorescence spectrum of the recombinan ANT(6) enzyme (Figure 2b).



	CONTIN	CDSSTR	SELCON3	$\overline{x} \pm \sigma$
% α-helix	48.2	45.8	47.7	47.2± 1.3
% β-sheets	10.3	14.5	11.7	12.2± 2.1
% Turns	13.5	15.6	15.8	15 ±1.3
%Random coil	28	24.9	27.3	26.7± 1.6

Figure 2 (a) Far-UV circular dichroism spectrum of ANT(6) in 5 mM Tris buffer of pH 7.5; (b) Fluorescence-emission spectrum of the ANT(6). The solid line represents the spectrum obtained upon excitation at 275 nm. The dotted line represents the contribution of tyrosines to the global spectrum.

The thermal stability of the enzyme was initially assessed by using CD spectroscopic analysis, in which the temperature dependence of the dichroism signal at λ =222 nm was monitored (Fig. 3). The curve following the thermal unfolding of the ANT(6) could be fitted to a simple thermodynamic unfolding model. The melting temperature of the recombinant ANT(6) was determined to be 56±0.2°C. The transition region was sharp, thus indicating that the protein existed initially as a compact, well-folded structure and that the unfolding reaction was highly cooperative.



Figure 3. Thermal denaturation profile of the ANT(6) enzyme. Melting temperature (T_m) was assessed using CD by following the temperature dependence of the dichroism signal at 222 nm.

The influence of divalent cations in the activity and stability of recombinant ANT(6)

The quaternary structure of the enzyme determined by sedimentation velocity experiments (Figure S1a) showed that the presence of Mg²⁺ strongly affected the aggregation state of the enzyme. The magnesium ion is by far the most frequently found metal ion cofactor in enzymatic systems. This is most likely related to the ability of Mg²⁺ to form stable complexes with ATP, which is normally associated with the metal ior when acting in a physiological environment. On the othe hand, Mn²⁺ is the required cofactor in many other transferase enzymes, such as glycosyltransferases of the Leloir patway.²⁷ Therefore, we decided to study the influence of Mg²⁺ or Mn² ion on the 2D-structure (Figure 4).

Figure 4. CD spectra at 25 °C of the ANT(6) enzyme at different concentrations of MgCl₂ and MnCl₂.

The ANT(6) showed a similar activity in the presences of Mn_{u,2} and MgCl₂ (0.07 and 0.08 U/mg, respectively) but was mu₁ more stable in the presence of Mg²⁺ (Figure 4). Concentratio \circ of MgCl₂ above 20mM promote a conformational change \circ f the protein that is reflected in the decrease of the peaks \approx 195 \approx 208 and \approx 222 nm, which corresponds to an increase in betasheet structure of about 7% but which maintains a relatively

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high degree of structure. However, manganese promotes a higher loss of the protein's secondary structure. Thus, from a concentration of 10 mM, the CD spectrum gives almost no signal indicating that the protein has lost its secondary structure and, therefore, its chirality. This effect could be due to the higher flexibility of Mn2+ ligand bonds in both length and angle, which could promote the conformational change of the recombinant protein.²⁸

ANT(6)-substrate specificity

The main kinetic parameters for ANT(6) from B. subtilis are displayed in Table 1..

Table 1.- Kinetic parameters of the ANT(6) from *B. subtilis*^[a]

Substrate	K _M	$k_{\rm cat}$	$k_{\rm cat}/K_{\rm M}$
	(M)	(s ⁻¹)	(s ⁻¹ ·M ⁻¹)
Streptomycin	3.8·10 ⁻⁵	0.03	$9.2 \cdot 10^2$
ATPMg	8.4·10 ⁻⁵	0.018	2.1·10 ²
Streptidine	6 10 ⁻⁴	0.0006	1.2

^[a]0.5mM of streptomycin, 2mM [ATPMgCl₂], pH 7.5.

To the best of our knowledge, the kinetic parameters of only three other ANTs have been described: the bifunctional enzyme ANT(3")-Ii/AAC(6')-IId from Serratia marcescens,²⁸ the ANT(2")-I from Klebsiella pneumoniae^{29,30} and the ANT(4') from S. aureus.^{31,32} The ANT(3")-li domain carried out the adenylation of streptomycin with a higher catalytic efficiency $(k_{cat}/K_{M} 3,210^{5})$ than that of the ANT(6) from *B. subtilis* (Table 1). This domain was highly specific for streptomycin; that is, it did not adenylate 4,5- or 4,6-aminoglycosides, thus showing a similar behaviour to that of the ANT(6). In the case of the ANT(2")-I/ANT(4'), its catalytic efficiency is similar to that of our enzyme but it does not recognize streptomycin as an aminoglycoside.

Figure 5. The effect of the magnesium free-ion in the adenylated activity of the streptomycin with the ANT(6)

In addition, we studied the effect of $MgCl_2$ concentration or the enzyme activity as a function of the relationship [MgCl₂]/[ATP]. The maximum activity was reached at a 17 m V concentration of free Mg^{2+} (Figure 5). Above this concentration, and with a free-magnesium concentration of 37 mM, neither increased activity nor an inhibitory effect was detected. The concentration of magnesium required to reach maximum activity was higher than that required to saturate the ATP. Some authors suggest that this may be because th€ enzyme requires two molecules of the divalent ion to carry out its activity: one that would bind to the ATP and the other that would bind directly to the enzyme¹⁵, which has been observed in the 3D-structure of ANT(4')/Kanamycin Complex (pdb;1kny)

We have already showed that streptidine is the minimum moiety of streptomycin that is recognized by the enzyme and that can be adenylated by the ANT(6) (Figure S5).²² In order to study the ANT(6) specificity, the enzyme activity was tested using different aminoglycosides, such as streptomycin neomycin, kanamycin, spectinomycin, ribostamycin, paromomycin, neamine and the 2-deoxystreptamine (the common moiety of the 4,5- and 4,6-aminoglycosides). The enzyme was highly specific for streptomycin (Table S1) and dic. not recognize 4,5-or 4,6-aminoglycosides (no activity wa: detected with kanamycin or ribostamycin). In fact, the enzyme only provides acceptor groups for hydrogen-bond interaction with aminoglycoside substrates. The abundance of acidic residues in the binding pocket of the enzyme can be readily explained by reference to the fact that aminoglycosides are invariably positively charged molecules. The other ANT enzymes described, such as ANT(4') and ANT(2"), accept different aminoglycosides as substrates, thus showing hi rates of substrate promiscuity^{31,32}. This differing behavior can be explained by reference to the fact that, in the former case, the natural substrates are 4,5 or 4,6-aminoglycosides, both o which share the 2-deoxystreptamine moiety (Figure 1).

We have also studied the specificity of the nucleoside triphosphate (ATP, GTP, CTP and UTP). The enzyme recognizes the nucleotides with similar affinities²⁵ but only showed activity with ATP/GTP (Table S2). According to these data ANT(6) shows a clear preference for nucleotide triphosphates incorporating purine aromatic systems. On the other har J, modifications of the inorganic fragment led to significant changes in activity (Table S2). Thus, replacement of a single oxygen atom by a methylene group (ATP versus AMPCPP) and the suppression of the terminal γ -phosphate group (ATP versus ADP, AMP) lead to a total loss of activity. Recent, results obtained by our research group have demonstrated that ANT(4') from S. aureus can employ inorganic triphosphate (P3) as a substrate to promote the regioselective phosphorylation of aminoglycosides.³³ However, in the case of the ANT(6), 🥠 reaction of phosphorylation with streptomycin/P3 w detected.

In vivo activity of ANT(6)

Streptomycin is powerful antibiotic with minimal inhibitory concentrations (MICs) of 5 µg/ml for E. coli BL21(DE3) strain. However, when the ANT(6) enzyme is present, it lost its antibiotic activity (MIC>200 µg/ml). On the other hand, streptidine and 2-deoxystreptamine (2-DOS) (common moieties of the streptomycin and 4,5- 4,6-aminoglycosides, respectively) did not show any antibiotic activity in the absence of ANT(6) (MIC >100 $\mu g/ml)$ (Table 2). In the case of 4,5-aminoglycosides, binding to neomycin was detected but no reaction was observed (Table S2).²⁵ As we have described, the streptidine is a substrate of the ANT(6), so when it is added together with streptomycin, it works as a "decoy acceptor".²² With these previous results in mind, we assayed the 2-DOS³⁴ as a "decoy acceptor" of the ANT(6) in the presence of streptomycin. Meanwhile, streptidine is able to reduce the MIC for streptomycine to 10 μ g/ml when ANT(6) is expressed, but 2-DOS did not show any affect (Table 2). This result could indicate that ANT(6) requires the presence of a guanidine group in the aminocyclitol, since it is so specific for streptomycin, unlike the other ANTs hitherto described. This was verified because AMP-2-DOS product was not detected by ESI-MS or HPLC.

Table 2.- *In vivo* activity of streptomycin and neomycin alone or in combination with streptidine against *E. coli* strains expressing or not the ANT(6).

Amynoglycoside	E. coli	MIC ^[a] <i>E. coli</i> (pET)	E. coli pET-aadk
Streptomycin ^[b]	5	5	>200
Streptidine	>100	>100	>200
Streptomycin+streptidine ^[b]	5	5	10
2-DOS ^[c]	>100	>100	>100
Streptomycin+2-DOS	2	100	100

 $^{[a]}$ MIC: Minimum Inhibitory Concentration, values (µg/mL).

^[b] Data taken from reference ^{22 [c]} 2-DOS (2-Deoxystreptamine)

Experimental

Materials.

pET-28b(+) vector was obtained from Novagen. Taq DNA polymerase was purchased from Ecogen. T4 DNA ligase was obtained from MBI Fermentas. Restriction enzymes *Eco*RI and *Hind*III were purchased from Boehringer Manheim. Isopropyl-L-thio- β -D-galactopyranoside (IPTG) was purchased from Applichem. *E. coli* competent cells BL21 (DE3) was purchased from Stratagene Co. (San Diego, CA). Kanamicyn, ATP and other aminoglycoside antibiotics were obtained from Sigma. Nickel-iminodiacetic acid (Ni²⁺⁻IDA) agarose was generous donated by ABT. *Bacillus subtilis* was obtained from CECT.

Cloning, expression and purification the ANT(6)

PCR amplification was performed in a 100 µL reaction mixture containing DNA (3 µL) of Bacillus subtilis as template, water (70.5 μL), buffer (100 mm Tris-HCL, 500 mm KCl, pH 8.0, 10 μL), MgCl₂ (2 mm), dNTPs (200 μm), primers (1 μm) ANT6-C⁺ (5'cgaagctttcactttactgagcaataaa3') and ANT6-Nt (5'gccgcgaattccatgcgaagtgagcaggaaat3') and Taq DNA polymerase (2.5 U). The reaction was subjected to 25 cycles of amplification. The cycle conditions were set as follows: denaturation at 94 °C for 1 min, annealing at 55°C for 1.0 min, and elongation at 72 °C for 1.5 min. The PCR product was introduces into the pGem-T easy vector The sequencing of the DNA insert was carried out with the promotors T7 and SP6. Then *aadK* insert was digested with *Eco*R and HindIII enzymes and inserted into the pET-28b(+) vector. that the resulting plasmid pET-ANT(6) was transformed into E. coli struction BL21 (DE3) which then showed the expected resistance pattern the streptomycin.

The selected clone was grown up on 100 mL of LB mediu. containing 26 µg/mL kanamycin at 37 °C with shaking. When the cell growth reached an optical density at 600 nm (O.D.600) of 0.5 the temperature was switched to 30 °C and the culture was induced with 0.5 mM IPTG. After 24 hours of induction the expression lev was analysed by SDS-PAGE using gels with 13% of polyacrylamide in the separation zone. The culture broth was centrifuged (5,000 x g 20 min, 4 °C), and the cells were suspending in Tris buffer (8 mL/g cells, 50 mm pH 8.0) and was added EDTA (50 mm, pH 8.2) and lysozyme (2 mg/g cells). The suspension was gently stirred at room temperature during 1 hour, and the suspension was kept at 4 °C overnight. The preparation was gently sonicated for 40 s and cooled down in ice (4 times) to decrease viscosity. DNase (10 μg/g cells) and MgCl₂ (0.95 μ g/mL of preparation) were added, and the mixture was refrigerated for 20 min. The mixture was then centrifuged for 30 min at 13,000 x g to separate the soluble proteins from the insoluble ones. To the soluble fraction 1% of streptomycin was added and stirred for 20 min at 4°C. Then the cel' free extract (CFE) was recovered by centrifugation.

1.5 mL of CFE was applied to 0.5 gr of Ni²⁺⁻IDA-agarose, previously equilibrated with phosphate buffer, pH 7.0. The proteins non specifically retained were washed with the same buffer. ANT(6) was eluted with the same buffer containing 0.5M of imidazole. Afte that, the solution was dialyzed against phosphate buffer 5mM, pH=7.0. After this step, the recombinant enzyme was found to re more than 90% pure as assessed by SDS-polyacrylamide g electrophoresis. Further purification of the recombinant ANT(6) war accomplished by size-exclusion chromatography on HiLoad 26/60 Superdex 200 PG column controlled using the AKTA-FPLC system The column was developed in 20 mM phosphate buffer pH 7.5 containing 0.15M NaCl at a constant rate of 1 ml/min. Fractions showing activity were pooled and dialyzed with water and concentrated by lyophilisation. Protein concentrations were estimated by the Bio-Rad protein assay method using bovine serven albumin as a standard.

Circular Dichroism spectroscopy.

Far-UV circular dichroism (CD) spectra were recorded in the wavelength range of 195-240 nm using a Jasco J-81(

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spectropolarimeter equipped with a constant temperature cell holder Jasco PTC423-S Peltier. The protein concentration was 5.3 μ M and the optical path length 0.1 cm. The contribution of the buffer was always subtracted. For each sample, three spectra were accumulated at a scan speed of 20 nm /min with a bandwidth of 0.2 nm and averaged automatically. All spectra were background-corrected, smoothed and transformed into mean residue ellipticity (θ) in the JASCO spectra manager. A value of 110 g/mol was used as a mean residue weight for ANT(6).The secondary structure of the protein was evaluated by computer fit of the dichroism spectrum into four simple components (α -helix, β -sheet, turns and random coil) using the CDPro software package containing three commonly used programs: SELCON3, CONTIN/LL and CDSSTR.²⁶

Ultracentrifugation.

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The quaternary structure of recombinant ANT(6) from *B. subtillis* was assessed using sedimentation equilibrium experiments. These experiments were performed at the Department of Chemical Physics of Biological Macromolecules of the IQFR-CSIC. The initial concentration of the protein used in these experiments was 0.96 mg/mL.

Fluorescence Experiments.

The experiments were conducted on Perkin Elmer LS50B luminescence spectrometer with the program FLWin. A quartz cell with a 1 cm path length in both the excitation (275nm and 295nm) and emission (295 to 400nm) directions was used in all the experiments that were performed at 25°C. The ANT(6') protein concentration was 30 μ M in phosphate buffer 5mM, pH=7.5.The spectres were normalized by origin 7.0 program.

Measurement of Enzyme activity.

The activity of the pure recombinant enzyme was follow by HPLC and the identity of the adenylated products were confirmed by ESI-MS. The formation of the product was followed by HPLC at 260nm (Vydac column eluted with TFA pH 3.0 buffer). Enzyme concentration was calculated using a molar absortion coefficient at 60400 M^{-1} cm⁻¹ at 280 nm. The reaction mixture contained 50 mM Hepes buffer, pH (6.5-8), nucleoside derivatives (3mM), aminoglicoside (0-2.5 mM), MgCl₂ (0-35mM) and the reaction were started by the addition of ANT(6) enzyme (0.5 μ M) at room temperature in a volume of 1 ml.

Minimal inhibitory concentrations (MIC)

The different strains were grown in 1 ml tubes and Mueller-Hinton broth to an optical density OD_{600} of 0.4 units, then ANT(6) was expressed by induction with 0.5 mM of IPTG until OD_{600} 0.6 when. Then, the desired concentrations of antibiotics were added from stock solutions. The samples were incubated at 37°C for 12 h, when the control culture (with no antibiotic) had an OD_{600} of 1.2-1.6. The OD_{600} of each sample was read and the MIC was taken as the lowest antibiotic concentration inhibiting bacterial growth greater than 90%.

Conclusions

In conclusion, we have described the overexpression and biochemical characterization of the ANT (6) from *B. subtilis*.

The results reported throughout this manuscript highlight the high specificity of the ANT(6) for streptomycin-ATP/GTP-Mg of an enzyme that is involved in bacterial resistance agains' aminoglycoside antibiotics.

From a medicinal chemistry perspective, the combination of the features present in the ANT(6) give the enzyme a narrow tolerance to chemical variations in the aminoglycoside/nucleotide, making it very useful in the design of non-inactivable derivatives in order to avoid bacterial resistance. Moreover, the design of new inhibitors or decov acceptors for this enzyme must considerer the minimum unit of streptidine that is essential, and the fact that only nucleotide derivatives of purine bases will be recognized.

Acknowledgements

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Notes and references

- [‡] The term "decoy acceptors" was introduced by M. Okayama, et all.²³ and N. B. Schwartz²⁴ to designate those compounds, that being substrates for a particular enzyme are able to inhibit a metabolic pathway, to distinguish them from inhibitors, which are compounds that block the enzyme without being modified by it.
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the aminoglycoside/nucleotide, making it very useful in the design of non-inactivable derivatives.