Poster Presentations

[MS5-P35] Binding of a G-quadruplex from *Streptococcus pneumoniae* to [Ru(phen)2dppz]²⁺ <u>Sarah P. Gurung</u>, James P. Hall, Christine J. Cardin, John A. Brazier,

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polypyridyl Ruthenium complexes have recently been shown to interact with a range of nucleic acid structures.[1-3] ([Ru(phenant hroline)2dipyrido[3,2- $a:2^{\prime},3^{\prime}c$]phenazine]²⁺) emits luminescence brightly in the presence of duplex DNA and acts as a "light switch" for G-quadruplexes.[3] A racemic mixture of Λ - and Δ - [Ru(phen)2dppz] was added to an in- house synthesised G-quadruplex forming se quence,d(GGGCTAATAGGGAGAGCAGGG ACGGG), which is predicted (via QuadFinder) to form within 14 strains of Streptococcus pneumonia but in varied locations. The nucleotide search tool, blastn, shows that Streptococcus pneumoniae and Streptococcus pseudopneumoniae are the only identified bacteria within which the particular G- rich sequence is always present. The binding of the complex to the G-quadruplex was studied under UV-Vis over a temperature range (20-90°C) both with/out dilute K where the quadruplex formed readily in K (100 mM).[4] The sequence complementary to the G-quadruplex, the i-motif, was also synthesised and its UV-Vis spectra showed stability, as indicated by the presence of protonated-cytosine base pairs, [5] under acidic conditions with a mean melting temperature $(T_{\rm o})$ of 46°C. Binding of the complex to the i-motif will be studied. Structures of these two sequences from Streptococcus pneumonia are to be examined with circular dichroism (CD) spectroscopy and their crystallisation attempts to be made with [Ru(phen)2dppz]² as well as other Ru-containing complexes, which, if positive with the capability to diffract well, could help understand the notion of i- motifs as possible therapeutic targets.

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