Analysis of New HIV-1 Inhibitors as Potential Antiviral Agents for HIV-2

Rowan Brothers¹, Daniel Kneller², Andres Wong-Sam², Johnson Agniswamy², Arun K. Ghosh³, Irene T. Weber¹,²
¹Department of Chemistry; ²Department of Biology, Georgia State University, Atlanta, GA
³Department of Chemistry and Department of Medicinal Chemistry, Purdue University, West Lafayette, IN

Human immunodeficiency virus type 2 infects approximately 2 million people worldwide, primarily those in or around West Africa. Due to its crucial role in the maturation of the virus, HIV-1 protease has been commonly used as a target for developing anti-retroviral drugs for the more prevalent HIV-1 infection. Hence, HIV-2 infections are currently being treated with HIV-1 protease inhibitors, often with decreased efficacy. Here, in order to evaluate potential new inhibitors, the HIV-2 protease is prepared for enzyme inhibition assays and structural studies by X-ray crystallography. Protein was prepared by bacterial overexpression of HIV-2 protease and purified using fast-protein liquid chromatography (FPLC) and high-pressure liquid chromatography (HPLC). Once purified this protein will be assayed with different potential inhibitors and crystallized for structural analysis to determine the key interactions between the inhibitor and the protease. New tight-binding inhibitors will have potential as antiviral agents for patients with HIV-2 infections.