Review Article **JPRR (2018) x:xx**

Journal of Pharmaceutical Research and Reviews (ISSN:2576-8417)

Lead Developer Angular and Effect of Force by Side Chain of Suicide Molecule in HIV AIDS Drug Discovery

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ABSTRACT

Angular to lead (A2L) accepted as lead generation has stage in early drug discovery where small molecule hits from a high throughput screen (HTS) has evaluated and undergo limited optimization to identify promising lead compounds. These lead compounds undergo more extensive optimization in a subsequent step of drug discovery. Angular imagination → Target validation (TV) → assay development → high-throughput screening → hit to lead (H2L) → lead optimization (LO) → preclinical drug development → clinical drug development.

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How to cite this article:
Rahul Hajare.Lead Developer Angular and Effect of Force by Side Chain of Suicide Molecule in HIV AIDS Drug Discovery. Journal of Pharmaceutical Research and Reviews, 2018; x:xx.

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Executive Statement

HIV has true evolutionary history of a homologous of sequences. It has mutations which produce molecule inactive [1,2]. It has some other property and impact on the fitness of the HIV carrier indicates how these analyses can be useful in the drug discovery process. HIV has a same organ under every variety of form and function. HIV has a variety of forms that can be changes by the sometime this science still holds secrete even today and selection removes variation. It has homologous proteins those differ in their specificities, specific activities, level of aggression or some other basic property. However homologous will have some residual similarity in their forms and function. HIV has a force that is neutral early stage with respect to natural selection; much of the extent sequence that is functionally important has experienced selective pressures in the beginning, governing gene and protein sequence. Mutation has the major source of variation [3,4]. Suicide inhibitor has picture to treat HIV infection in mutational stages. HIV has been separated due to difficult to understand and study that deals with the study of identification, structure, rate of growth and productivity of HIV virus in host. HIV has heterogeneous group as well as homogeneous group of several distinct classes of living sub type of structure. HIV having different types of hetero atom along with fused heterocyclic compound with known and unknown odd mechanics; it has quantum mechanics that has invisible in 3D function [5, 6]. Physician remark for outcome of therapy for ART reveal the information about patient's related factors including medication adherence. It has brought some useful innovations to the study of treatment. It was felt that it should not be separate drug discovery standard for reduce the dose frequency and reduce adverse effect and short time treatment. Suicide inhibitors has natural history of drug discovery could illustrate the complexity in selecting a baseline of molecule. Drug discovery is a relatively slow growing chemical process with an incubation period said to range from a few year to a life time. Suicide drug discovery has an extra benefit in any case of side effect is detected, it should be treated thoroughly.

Introduction

The force is the sum of all the forces between two neighbouring molecules. The forces result from the actions of the kinetic energy of atoms and the slight positive and negative electrical charges on different parts of a molecule that affect neighbours and make absolute that may be produce effect [7, 8]. The main categories of intermolecular forces are bond interaction, interaction makes molecule stable and stable molecule always produce significant results in HIV. Hydrogen bonding is considered a form of dipole-dipole interaction, and so contributes to the net intermolecular force. In contrast, intramolecular force is the sum of the forces that act within a molecule between its atoms. The intermolecular force is measured indirectly using measurements of various properties, including volume, temperature, pressure, and viscosity. This effect has known as suicide effect. Suicide inhibitors come in picture to treat HIV in stable as well as steady effect. In suicide inhibition, also accepted as suicide inactivation is quantum-based inhibition, is an irreversible form of enzyme inhibition that occurs when an enzyme binds a substrate analogue and forms an irreversible complex with it through a covalent bond (C- H) during the HIV reaction [9,10]. The inhibitor binds to the active site where it has compared modified by the enzyme to produce a reactive group that reacts irreversibly to form a stable inhibitor-enzyme linkage. This usually uses a prosthetic group or a coenzyme, forming electrophilic alpha (E^{+}) and beta unsaturated carbonyl compounds and imines (stable electrophilic and nucleophilic intermediate attachments) [11, 12]. Suicide inhibitors has used in what is called "rational drug process" where the aim is to create a novel substrate, based on already known mechanics and substrates. The main

goal of this approach is to create substrates that has unreactive until within that enzyme's active site to reach at proper HIV functional group and at the same time being specific that can produce high efficacy. Drugs based on this approach have the advantage of very few resulting side effects [13,14].

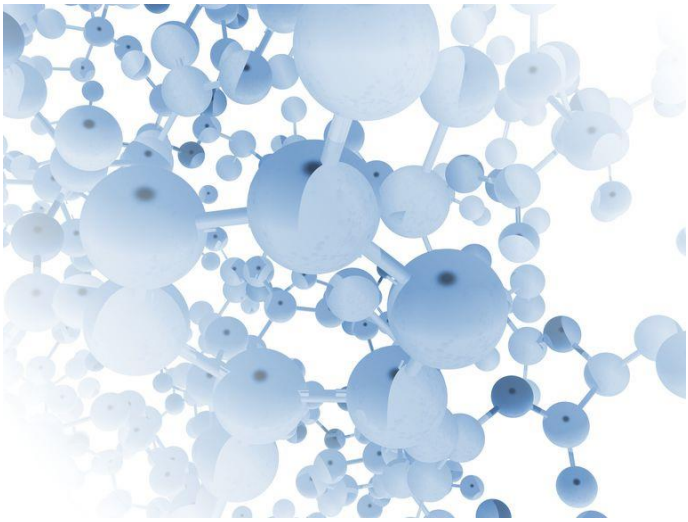

Fig 1 Intermolecular forces of molecule are those that occur between suicide molecules and HIV

Conflicts of interest

None

Funding

None

Acknowledgment

This study has been guided under the supervision and guidance of Respected Dr. Ramesh Paranjape,, Founder National AIDS Research Institute India. I express my deep gratitude towards Respected Sir, for motivation and being great knowledge source for this work.

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