Molecular docking based screening of Listeriolysin-O for improved inhibitors

Sara Ghafari1, Matin Komeilian1, Mohaddese sadat Hashemi1, Sareh Oushani1, Garshasb Rigi2, Behnam Rashidieh1, Kamran Yarahmadi1, Fatemeh Khoddam1*

1Vira Vigene research institute, Tehran, Iran; 2Department of Biology, Faculty of Science, Behbahan Khatam Alanbia University of Technology, Behbahan, Iran; *Corresponding author: Fatemeh Khoddam - Email: elnazkhodam@yahoo.com

Received January 17, 2016; Revised April 23, 2016; Accepted April 23, 2016; Published May 31, 2017

Abstract:
Listeriolysine-O (LLO) is a 50KDa protein responsible for Listeria monocytogenes pathogenicity. The structure of LLO (PDB ID: 4CDB) with domains D1 to D4 is known. Therefore, it is of interest to identify conserved regions among LLO variants for destabilizing oligomerization (50 mer complex) of its monomers using appropriate inhibitors. Therefore, it is of interest to identify suitable inhibitors for inhibiting LLO. Previous reports suggest the use of flavanoids like compounds for inhibiting LLO. Our interest is to identify improved compounds to achieve its fatality especially in pregnant women following to L. monocytogenes' toxin proteins so called: Listeriolysine-O (LLO) is contributing to the pathogenicity of the organism and enables it to achieve its fatality especially in pregnant women following to invade the tissues of the host thus it is known as a virulence factor [4,5]. Particularly the hallmark of this factor are to be non-enzymatic, cytolytic, thiol-activated, cholesterol-dependent, pore-forming toxin and most notably it remains in an active form even after bacteria's death [6]. It also can induce cytolysis in infected host cells even in low concentration of 5 ng/ml [7]. Furthermore, LLO mediated cell death is proceeded through cytolysis or apoptosis [8]. Apoptotic event, which is mediated by LLO, can induce two pathways: mediating by activation of caspase-3 and caspase-6, another pathway dependent on LLO but independent to caspase [9]. First pathway in apoptotic T-cells have been described. Activation of mechanism by 3 surface receptors starts. Activated T-cells have obtained FAS L as ligand receptors. Attachment of LLO as a ligand to FAS L leads to activation of death domain, which attaches to the outer domain. Death domain also plays a pivotal role in connecting caspase as an adaptor, so this interaction can activate pro-caspase. This factor realizes caspases-3 that can inactivate DNA. All these factors end in segmenting DNA in the host cell [10].

Keywords: Listeria monocytogenes, Listeriolysine-O, Molecular docking, Drug discovery

Background:
Listeria monocytogenes is known for its virulence among the most famous species of Listeria [1]. Nevertheless this gram-positive bacterium is a facultative intracellular pathogen; the disease might be occurred due to germ entrance through contaminated foods or beverages into a healthy body [2]. While this bacterium is to some extent resistant to Gastric acid and Bile salts [3], one of the L.monocytogenes's toxin proteins so called: Listeriolysine-O (LLO) is contributing to the pathogenicity of the organism and enables it to achieve its fatality especially in pregnant women following to invade the tissues of the host thus it is known as a virulence factor [4,5]. Particularly the hallmark of this factor are to be non-enzymatic, cytolytic, thiol-activated, cholesterol-dependent, pore-forming toxin and most notably it remains in an active form even after bacteria’s death [6]. It also can induce cytolysis in infected host cells even in low concentration of 5 ng/ml [7]. Furthermore, LLO mediated cell death is proceeded through cytolysis or apoptosis [8]. Apoptotic event, which is mediated by LLO, can induce two pathways: mediating by activation of caspase-3 and caspase-6, another pathway dependent on LLO but independent to caspase [9]. First pathway in apoptotic T-cells have been described. Activation of mechanism by 3 surface receptors starts. Activated T-cells have obtained FAS L as ligand receptors. Attachment of LLO as a ligand to FAS L leads to activation of death domain, which attaches to the outer domain. Death domain also plays a pivotal role in connecting caspase as an adaptor, so this interaction can activate pro-caspase. This factor realizes caspases-3 that can inactivate DNA. All these factors end in segmenting DNA in the host cell [10].

Köster et al have revealed Crystal structure of LLO in in 2014. According to the crystal structure, it consists of four distinct domains, which have been called D1 to D4 and each of them, playing a different role in LLO functionally. This study was conducted to intend to illustrate some sequences of LLO, which are more conserved in D1 to D4. Describing these sequences can be of a great importance for modeling an inhibitor for inhibiting oligomerization of LLO monomers [11]. Two conserved sequences in D1 can be noted. Firstly, the sequence contained 25 amino acids, which are significant for LLO function. Additionally, LLO has...
another important region that is known as a PEST-like sequence (P: proline, E: glutamine, S: serine, T: threonine). This region is essential for virulence and L. monocytogenes’s in-vivo function [12]. Moreover, 91-99 sequences of D2 play a key role in immunogenic and identified by CD+8 T-cells but a highly conserved motif structure in D4 consists of 11 residues and makes contribution to cytotoxicity of L. monocytogenes. LLO is able to connect to the intestinal epithelial by Internalin protein. Also, LLO is one of those factors, which released by infected cell to other cells. Pore forming is a mechanism that causes transition and during this procedure a hole is created in host cell resulted in cytolysis in the host cells and finally leads to cell death [13].

During past these years, some drug targets were introduced with bactericidal mechanism: Yasuhiro Gotoh et al. (2010), worked on specific inhibitor against two component signal transduction systems (TCSs) which could reduce virulence of bacteria with inhibiting the sensory domains of the sensors blocking the quorum sensing system [14]. One year later, Mikael Mansjo along with Jorgen Johansson introduced FMN riboswitch as a novel drug target for antibacterial substances. They investigated how flavine analog, roseoflavin, affected the growth and infectivity of L.monocytogenes at a very low concentration. Interestingly, their results admitted that roseoflavin enhances L.monocytogenes virulence in mechanism independent of the FMN riboswitch [15]. Wang J et al. also showed the detection of LLO native inhibitors with contrasting activity by using hemolysis test [16] but the inhibitors should be improved. LLO is pore forming toxin (PFTs) and its monomers oligomerize into ring of 50 monomers. Accordingly this virtual experiment, such as our other studies in this field [17, 18, 19], is conducted to aim for modeling an essential inhibitor for prohibiting oligomerization of LLO monomers, which causes to induce inhibiting of oligomer formation and prevent pore forming of LLO [20, 21, 22, 23].

Figure 1: A: The structure of LLO, it consists of four distinct domains, which have been called D1 to D4 and each of them, playing a different role in LLO functionally. B: The modified structure (RD-1) in contact with D2 domain of LLO. C: The unbound RD-1 structure. D: The overall pharmacological properties of RD-1. E: The ligand map of RD-1. F: The structure of Hit#1. G: The structure of Hit#2. H: The structure of Hit#3. I: The structure of Hit#4. J: The structure of Hit#5
Methodology

Protein and ligand structures

The X-ray crystal structure of 488 amino acids polymer of *L. monocytogenes* toxin (LLO) obtained from protein data bank (www.rcsb.org/pdb/home/home.do) with pdb access code of 4CDB [11]. The provided structure was a pore containing up to 50 monomers with a diameter of 300 Å. Considering the aim of this study that is to inhibit monomer interactions, the chain A of this Alpha-helical Pore-forming toxin was separated as the target for drug binding. Besides, to provide biological conditions simulation, the monomer structure was solvated in a water box with the distance of 2 Å. Adding Na or Cl ions then neutralized the system. The solvation process was carried out by GROMACS 4.5.6 simulation software. The zinc database (http://zinc.docking.org/) was used as the ligand database for virtual screening [24]. A subset from drug-like category obtained and used as virtual screening library. Virtual screening was performed among 200,000 ligands and top successive hits were selected for rational drug design purpose.

Virtual screening:

PyRX software [25] was used as the virtual screening software. PyRX includes Autodock vina [26] with a Lamarckian genetic algorithm as scoring algorithm.

Pharmacokinetic analysis and rational drug design:

FAF Drugs 3 web server [27] was used to analyze the absorption, distribution, and metabolism properties of top 10 virtual screening hits. Moreover, the toxicity properties were analyzed using the PROTOX web server [28]. New ligands were designed based on the structure of top hits retrieved from virtual screening process. To achieve this goal, hyperchem software was used. New rationally designed ligands were then analyzed regarding ADME and toxicity to reach optimal scores.

Results and discussion:

Among 200,000 drug-like ligands, the 5 highest binding affinity hits were selected for the further evaluation. These top 5 poses, which indicated more negative binding affinity, were examined for pharmacokinetic analysis and rational drug design purpose. The selected hits reached the binding affinity equal to -9.6 for hit 1, -9.4 for hit 2, -9.1, -8.8 and -8.6 for hit 3, 4 and 5 respectively. Based on Lipinski rule of 5 we analyzed the hits regarding pharmacological properties. In table 1 the pharmacological properties of these top hits are presented. To reach the best inhibitor, we focused on PEST sequence in D1 of LLO structure, which is close to the N-Terminus. This sequence plays a key role in LLO dependent bacterial virulence. Hit 1 and hit 2 are very similar in structure just with one hydroxyl group differentiation. Other hits are very small so they seemed to be not suitable for being candidate as this limitation prevented protein-protein interaction correctly. For acquiring the specific binding affinity we tried to design specific ligands, which could tightly bind to D1 domain. To do this, we chose hit 1 because of high binding affinity and appropriate molecular weight. Thus, initially we endeavor to modify hit 1 to reach its optimal pharmacological properties. Based on FAFDrugs3 output, there are some errors in rotatable bonds and LogP. So based on hit 1, RD-1 was rationally designed and to decrease LogP, one carbonyl was added to the primary structure. Also in the base structure, there were problems in rotatable bonds. Changing rotatable bonds directly affects the binding affinity. Several refinement steps were applied in the base structure to decrease its rotatable bonds in a manner that the binding affinity remains still high. The propane group was removed from the structure end to implement this change and also five double bonds were added to solve the rotatable bonds problem. These structurally modifications and substitutions were changed the pharmacophore model seriously but the binding affinity still was -9.2. The new structure was checked again and still the same error was exits. To solve this problem, we have added a cyclopentane substitute and nitrogen was added in center of RD-1. These refinements were not significant effected other parameter such as complexity and tPSA. After these changes, RD-1 finally passed FAFDrugs successfully with the binding affinity equal to -9.1. The final structure of RD-1 in contact with LLO following its ligand map is depicted in Figure 1.

Table 1: The pharmacological properties of the top hits

| Ligand No | Compound | MB | HBdonar | HBa | Solubility Mg/l | Rotatable Bond | Rigid Bond | Sterocentres | Binding affinity |
|-----------|----------|----|---------|-----|----------------|----------------|------------|--------------|-----------------|
| 1         | 2-methyl-octadecane-1,3,4-triol | 316.52 | 3 | 3 | 2816.98 | 16 | 0 | 3 | -9.6 |
| 2         | 2-methyl-octadecane-1,3 -diol | 300.52 | 2 | 2 | 1422.18 | 16 | 0 | 2 | -9.4 |
| 3         | 2-ethyl-hexane 1,3-diol | 146.23 | 2 | 2 | 41706.33 | 5 | 0 | 2 | -9.1 |
| 4         | heptane 1,2,3,4,5,6,7-heptanol | 212.20 | 7 | 7 | 142623.86 | 6 | 0 | 5 | -8.8 |
| 5         | 3-hydroxymethyl-cyclohexane-1,2-diol | 146.18 | 3 | 6 | 75923.44 | 1 | 6 | 3 | -8.1 |
| RD-1      | 1-(4-Cyclopent-3-enyl-6, 7-dihydroxy-8-hydroxymethyl-nona-2, 8-dienylideneamino)-penta-1,4-dien-3-one | 345.43 | 3 | 5 | 41573.02 | 11 | 11 | 4 | -9.1 |

ISSN 0973-2063 (online) 0973-8894 (print)
Bioinformation 13(5): 160-163 (2017)
In order to calculate the LD50 and probable nonspecific targets, PROTOX webserver was used. PROTOX prediction indicated LD50 of RD-1 is 1170 mg/kg with toxicity class 4. This performance operated in average similarity 80.26% and prediction accuracy of 78.26%. Interestingly no toxicity target (nonspecific) was found by PROTOX as well. As FAFDrugs and PROTOX indicated, the RD-1 is a lead compound and can serve as a new drug to inhibit LLO.

Conclusion:
Identification of an LLO inhibitor to destabilize its 50-mer oligomerization is of interest. We describe the identification, modification and design of an LLO inhibitor named RD-1 with fitting properties for further consideration. It should be noted that further in vitro studies are needed to confirm binding and inhibition of LLO.

References:
[1] Liu D, J Microbiol Methods. 2007 71:2 [PMID: 17884210]
[2] Kathariou S, J Food Prot. 2002 65:11 [PMID: 12430709]
[3] Chung HS, Int J Food Microbiol. 1999 47: 1 [PMID: 10357270]
[4] Portnoy, D. A., & Paterson, Y. Infect Immun. 1992 60:4 [PMID: 1312514]
[5] Schnupf P et al. Infect Immun. 2007 75:11 [PMID: 17682039]
[6] Geoffroy C, Infect Immun. 1987 55:7 [PMID: 3110067]
[7] Jacobus T, Mol Microbiol. 1998 28:6 [PMID: 9680200]
[8] Carrero J. A et al. J Exp Med. 2004 200:4 [PMID: 15302900]
[9] Martinon F & Tschopp J, Cell Death Differ. 2007 14:1 [PMID: 16977329]
[10] Dudek NL, Diabetes 2006 55:9 [PMID: 16936188]
[11] Koster S, Nat Commun. 2014 5:3690 [PMID: 24751541]
[12] Decatur AL, Science 2000 290:5493 [PMID: 11062133]
[13] Pfeffer K, Cell 1993 73:3 [PMID: 8387893]
[14] Gotth J, J Antibiot. 2010 63:3 [PMID: 20111065]
[15] Mansjö M & Johansson J, RNA Biol. 2011 8:4 [PMID: 21596602]
[16] Wang J et al. Sci Rep. 2015 9:5 [PMID: 25749652]
[17] Abazari D et al. Bioinformation 2015 11: 5 [PMID: 26124568]
[18] Rigi G et al. Bioinformation 2017 13:2
[19] Rigi G et al. Journal of Paramedical Sciences 2017 8:2
[20] Carter AB, Am J Respir Cell Mol Biol. 1998 18:3 [PMID: 9490656]
[21] Schuerch D W, Proc Natl Acad Sci. 2005 102:35 [PMID: 16105950]
[22] Rashidieh B et al. Bioinformation. 2015 11:8 [PMID: 2640917]
[23] Rashidieh B et al. Bioinformation. 2015 11:11 [PMID: 26912950]
[24] Irwin J J & Shoichet BK, J ChemInf Model. 2005 45: 177 [PMID: 15667143]
[25] Dallakyan S & Olson AJ, Mol Biol. 2015 1263: 243 [PMID: 25618350]
[26] Trott OJ, Comput Chem. 2010 31: 455 [PMID: 19499576]
[27] Lagorce D et al. BMC Bioinformatics 2008 9: 396 [PMID: 18816385]
[28] Drwal M et al. Nucleic Acids Res. 2014 42: 53 [PMID: 24838562]