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http://hdl.handle.net/2440/105974
A high-throughput screening assay for eukaryotic elongation factor 2 kinase inhibitors

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Received 25 February 2016; received in revised form 21 March 2016; accepted 7 April 2016

\textbf{KEY WORDS}

eEF2K; Inhibitors; High-throughput screening; Luminescence; MH-1 peptide

\textbf{Abstract} Eukaryotic elongation factor 2 kinase (eEF2K) inhibitors may aid in the development of new therapeutic agents to combat cancer. Purified human eEF2K was obtained from an Escherichia coli expression system and a luminescence-based high-throughput screening (HTS) assay was developed using MH-1 peptide as the substrate. The luminescent readouts correlated with the amount of adenosine triphosphate remaining in the kinase reaction. This method was applied to a large-scale screening campaign against a diverse compound library and subsequent confirmation studies. Nine initial hits showing inhibitory activities on eEF2K were identified from 56,000 synthetic compounds during the HTS campaign, of which, five were chosen to test their effects in cancer cell lines.

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1. Introduction

Protein is the most important structural and functional macromolecule in the cell, therefore, the process of translation (also called protein synthesis), which translates a messenger RNA (mRNA) into a protein, is essential in this regard. Protein synthesis is usually divided into three main stages: initiation, elongation and termination. The initial stage involves a variety of translation initiation factors that work coordinately to provide stringent regulatory mechanisms of translation. The elongation stage consumes a large amount of energy and amino acids within cells to synthesize new polypeptide chain, which will undergo further post-translational modification and fold into correct quaternary structure to function in the cell. Indeed, more than 99% of the energy and nutrients used in protein synthesis are consumed during elongation. Studies on translation termination mainly focus on abnormal signals and their effects on mRNA degradation while the understanding of other aspects of this step is rather limited.

Human eukaryotic elongation factor 2 (eEF2) is a member of the GTP-binding translation factor family that promotes the movement of ribosome along mRNA from one codon to the next. During the elongation stage, eEF2 facilitates the translocation of peptidyl-tRNA from ribosome A site to P site. eEF2 can be phosphorylated and thus inactivated by human eEF2K. The phosphorylation of eEF2 plays an important role in the regulation of protein synthesis. It was widely recognized that under stress conditions such as nutrient starvation, hypoxia or acidosis, the activity of eEF2K in tumor cells was upregulated which allows tumor cells to adapt and survive in such an adverse microenvironment. Thus, eEF2K has been proposed as a potential target for cancer therapy.

Among the human kinome, the majority (90%) of protein serine/threonine and tyrosine kinases share a similar architecture within their catalytic domain, so they are classified as conventional protein kinases (CPKs). The remainder (about 10%), which displays little sequence homology with CPKs, are named as atypical protein kinases (APKs). APKs and CPKs lack sequence identity, but share a related catalytic core as well as a remarkably similar N-terminal lobe that predominantly folds into a curved sheet and contains the phosphate binding loop (P-loop). The interlobe cleft serves as ATP binding pocket and possesses conserved key catalytic residues. Despite having similar ATP binding pockets, the spatial location of their conserved region GXGXXG is different. eEF2K is such an atypical Ser/Thr-protein kinase that it was also known as ‘Ca²⁺/CaM-kinase III’. Human eEF2K is composed of 725 amino acids.

It was well established that phosphorylation at Thr56 inactivates eEF2 thereby regulating protein synthesis at the elongation step of translation. This mechanism appears crucial for cancer cell survival as it reduces energy and amino acid consumption to help them live through adverse conditions. This point of view is supported by studies on cellular microenvironment and metabolic stress with breast cancer cells. Since eEF2K is neither a conventional protein kinase nor has other known substrates, blockage of its action does not significantly affect normal biological processes. eEF2K inhibitors can become new drug candidates for cancer therapy.

It was thought that an eEF2 peptide (amino acid sequence: SARGAGTRFTDTRKDE) containing the phosphorylation site was the substrate of eEF2K. When tested in vitro, however, it did not work efficiently. Scientists subsequently found two protein kinases MHCK (myosin heavy chain kinase from Dictyostelium discoideum) A and B, whose structures had considerable similarity to eEF2K, could phosphorylate a 16 amino acid peptide MH-1 (amino acid sequence: RKKFGESEKTKTEF) in vitro, an action that can be replicated by eEF2K. Thus, MH-1 peptide has been used as a surrogate substrate for eEF2K.

In this study, we developed and validated a luminescence-based high-throughput screening (HTS) assay for the identification of eEF2K inhibitors by use of MH-1 peptide. Traditional method to detect the catalytic activities of protein kinase is to quantify the incorporation of radioactive γ-phosphate form [γ-32P] adenosine triphosphate (ATP) into a peptide or protein substrate by using a scintillation counter. This is impracticable for HTS application. Immunoblotting assay can also be employed in screening, but the sample preparation is complex and costs time and money: it is therefore not suitable for HTS and could be readily replaced by fluorescence and luminescence approaches. MH-1 peptide substrates have been commercialized. For instance, ADP-Glo™ luminescent kinase assay kit can eliminate redundant ATP in the assay system and transform ADP into ATP effectively. This became the key technology used in this study.

2. Materials and methods

2.1. Reagents

A484954, eEF2K enzyme produced and purified from Escherichia coli according to the method previously described and calmodulin were provided by Professor C. Proud at South Australia Health and Medical Research Institute. MH-1 peptide was purchased from China Peptides Co., Ltd. (Shanghai, China). EDTA, EGTA, CaCl₂ and MgCl₂ were bought from Shanghai Chemical Reagents Co., Ltd. (Shanghai, China). 3-(N-Morpholino)propanesulfonic acid (MOPS), DMSO and β-mercaptoethanol (β-ME) were obtained from Sigma (St Louis, MO, USA). The assay plates were the products of PerkinElmer (Boston, MA, USA). ADP-Glo™ luminescent kinase assay kit was procured from Promega (Wisconsin, MA, USA). The anti-eEF2 and anti-phospho-eEF2 antibodies were purchased from Cell Signaling Technologies (Boston, MA, USA). L15 medium, fetal bovine serum, penicillin, streptomycin, sodium pyruvate, genetin and 0.5% trypsin–EDTA were bought from Life Technologies (Carlsbad, CA, USA). Cancer cell lines used in this study were obtained from ATCC (Manassas, VA, USA).

2.2. Cell culture

MDA-MB-453 cells were maintained in L15 medium containing 10% fetal bovine serum (FBS), 100 units/mL penicillin/streptomycin, 1 mmol/L sodium pyruvate and 0.5 mg/mL gentamicin without CO₂. H1299 cells and HCT116 cells were maintained in RPMI-1640 medium and McCoy’s 5a medium containing 10% FBS, respectively. 100 U/mL penicillin/streptomycin, 1 mmol/L sodium pyruvate and 0.5 mg/mL genetin in a 5% CO₂ incubator.

2.3. Compound library

The compound library used for the screening of eEF2K inhibitors is consisted of 56,000 pure synthetic compounds. The stock was
presolubilized in 100% DMSO prior to application in the HTS campaign with an average concentration of 1 mg/mL for each compound. The purity (95% minimum) of all compounds was documented and randomly verified by the Chinese National Compound Library.

2.4. Luminescence-based HTS campaign

Before the primary screening, assays were performed manually in solid white 384 square well plates (BD Bioscience, NJ, USA). Serial dilutions of each reagent (enzyme, ATP and unlabeled peptide MH-1) were made to determine the optimal experimental conditions, which led to the selection of 6 ng/μL and 100 μmol/L for eEF2K and MH-1, respectively. In detail, we selected a 10 μL reaction volume. 4 μL of assay mixture 2 (MH-1 and ATP diluted in H2O) was first dispensed by Bravo liquid handler (Agilent, CA, USA) to an assay plate. 2 μL of compounds (1 mg/mL) were diluted with 18 μL 1× kinase buffer in 96 square well plates, then transferred (2 μL) to the 384 square well plates as described above. 4 μL of assay mixture 1 (eEF2K, β-ME and calmodulin diluted in 1× kinase buffer) was then dispensed to each well of the 384 square well plates. Enzyme was first mixed with compounds to initiate the reaction prior to incubation at room temperature (RT) for 2 h. The reaction was stopped by 10 μL of ADP-Glo™ luminescent kinase assay kit followed by incubation at RT for at least 40 min. At last, 10 μL of kinase detection reagent was introduced and incubated at RT for 30 min. Luminescence was measured with an EnVision plate reader and the emission filter used was luminescence 700 (PerkinElmer).

Hits identified from the primary screening were hand-picked (5 mg/mL in DMSO) and serially diluted (1:3) seven times to give a total of eight different concentrations. Each compound was tested in duplicate and the dose-response characteristics were analyzed using GraphPad Prism software (GraphPad, San Diego, CA, USA) for nonlinear regression analysis.

2.5. Western blot

Cells were seeded in 24-well plates at 2 × 10^5 cells/well. Twenty-four hours later, they were treated with A484954 and hit compounds for 6 h, rinsed once with Dulbecco’s phosphate-buffered saline (D-PBS) and harvested in 2× SDS-PAGE loading buffer (TakaRa, Dalian, China). Cell lysates were incubated at 95 °C for 10 min and then resolved in Criterion XT 10% gels (Bio-Rad, Hercules, CA, USA) for at least 60 min. Proteins were transferred to an Immobilon FL PVDF membrane (Millipore, Merck KGaA, Darmstadt, Germany) in 1× transfer buffer for 40 min. The membranes were rocked gently in blocking buffer (5% skimmed milk) and probed with primary antibodies to phospho-eEF2 at Thr56 and eEF2 at 4°C overnight. The antibody against β-actin was from Life Technologies. Detection of primary antibodies was performed using Alexa Fluor 680 against rabbit secondary antibodies from Invitrogen. Image acquisition and
analysis were performed using the ChemiDoc™ MP System (Bio-Rad).

2.6. Statistical analysis

All results were analyzed by using Prism 5 (GraphPad). $Z'$ factor is a simple statistical parameter for HTS assay, whereas signal-to-background ratio ($S/B$) is to assess assay quality. Data variation is analyzed by standard deviation (SD) or coefficient of variation (CV) $^{16}$. The statistical characteristics and percentage inhibition in this study were calculated as below.

\[
Z' = 1 - 3 \times \frac{SD_{PC} + SD_{NC}}{|Mean_{PC} - Mean_{NC}|}
\]

(1)

\[
S/B = \frac{\text{Mean signal}}{\text{Mean background}}
\]

(2)

\[
CV(\%) = \frac{SD}{Mean} \times 100
\]

(3)

\[
\text{Inhibition(\%)} = \left(1 - \frac{\text{Signal}_{\text{test}} - \text{Signal}_{\text{NC}}}{\text{Signal}_{\text{PC}} - \text{Signal}_{\text{NC}}}\right) \times 100
\]

(4)

where PC represents positive control wells; NC represents negative control wells; SD represents standard deviation, M represents mean.

3. Results

3.1. Assay optimization

A number of assay parameters that may affect signals evoked by kinase reaction and background noises were studied. In order to ascertain the optimal incubation time, we detected the final luminescence signal at different times after initiation of the assay.

Figure 3 Results of high-throughput screening campaign against eEF2K. Distribution of potential hits identified from 56,000 small molecule compounds is expressed as percentage inhibition of each sample compared with the positive control (A484954).

<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Chemical formula</th>
<th>Molecular weight</th>
<th>IC$_{50}$ (µmol/L)</th>
</tr>
</thead>
<tbody>
<tr>
<td>WNN0017-C003</td>
<td><img src="image" alt="Structure" /></td>
<td>C$<em>{23}$H$</em>{34}$BrNO</td>
<td>410.35</td>
<td>32.3</td>
</tr>
<tr>
<td>WNN0048-A002</td>
<td><img src="image" alt="Structure" /></td>
<td>C$<em>{27}$H$</em>{40}$N$^+$ Br$^-$</td>
<td>458.52</td>
<td>13.27</td>
</tr>
<tr>
<td>WNN0532-H011</td>
<td><img src="image" alt="Structure" /></td>
<td>C$<em>{19}$H$</em>{21}$F$_3$N$_2$O</td>
<td>345.39</td>
<td>0.032</td>
</tr>
<tr>
<td>WNN0572-A011</td>
<td><img src="image" alt="Structure" /></td>
<td>C$<em>{15}$H$</em>{12}$Cl$_2$O$_4$S$_2$</td>
<td>391.29</td>
<td>0.069</td>
</tr>
<tr>
<td>WNN0593-D007</td>
<td><img src="image" alt="Structure" /></td>
<td>C$<em>{19}$H$</em>{19}$NO</td>
<td>277.36</td>
<td>0.23</td>
</tr>
</tbody>
</table>
with appropriate enzyme and superfluous substrate. Since a reaction time of 2 h gave a good signal window it was set as an optimized incubation period (Fig. 1A). Considering that there is a time postponement between first and last batch of samples during the HTS process and kinase at RT may be unstable, we diluted the enzyme (initial concentration was 0.05 mg/mL) by 1:2–1:7 different concentration gradients and placed them at RT for 0, 2 and 4 h, respectively, before adding to the assay plate to activate the reaction. In this condition, according to the luminescence signal shown in Fig. 1B, 0.00625 mg/mL (or 6 ng/reaction. In this condition, according to the luminescence signal shown in Fig. 1B, 0.00625 mg/mL (or 6 ng/reaction) enzyme (initial concentration was 0.05 mg/mL) by 1:2–1:7 different concentrations and placed them at RT for 0, 2 and 4 h, respectively, before adding to the assay plate to activate the reaction. In this condition, according to the luminescence signal shown in Fig. 1B, 0.00625 mg/mL (or 6 ng/µL) enzyme displayed acceptable response which was thus set as an optimal concentration. Other reagents such as β-ME, ATP and calmodulin were introduced according to the amounts described above. Finally, DMSO (blank) and compound A484954 (positive control) were employed to calculate Z’ factor (0.73), S/B ratio (15.78) and CV (%7) values (Fig. 1C), which suggest that our assay system is of high quality and suitable for HTS.

After optimizing the assay conditions, we studied concentration-dependent response features of A484954. An IC_{50} value of 0.23 µmol/L, consistent with that reported in the literature (0.28 µmol/L), was obtained (Fig. 2).

3.2. HTS campaign

We applied 56,000 compounds from the Chinese National Compound Library to a large-scale random screening against eEF2K. Distribution of inhibition is illustrated in Fig. 3. Only nine initial hits showing inhibition over 25% were identified. They were hand-picked and rescreened using the same assay in duplicates. Five compounds (WNN0017-C003, WNN0048-A002, WNN0532-H011, WNN0572-A011 and WNN0593-D007) displaying consistent inhibition on eEF2K (>30%) were confirmed and further studied for their concentration–response characteristics (Table 1) and effects on cancer cell lines.

3.3. Effects on cells

A484954 is a highly selective eEF2K inhibitor with an IC_{50} value of 0.28 µmol/L in an enzymatic assay while having little activity against a wide panel of other serine/threonine and tyrosine kinases. Our Western blot data (Fig. 4) demonstrate that A484954 was capable of inhibiting eEF2 phosphorylation in MDA-MB-453, H1299 and HCT116 cells. Due to limited compound availability, we selected WNN0048-A002 (purity >95%) as a representative to investigate its activity in MDA-MB-453 cells. WNN0017-C003 was used as a negative control. Surprisingly, both compounds increased the phosphorylation of eEF2 (p-eEF2) in the cell and the total protein level of eEF2 was not affected by either A484954 or hit compound treatment.

4. Discussion

In this paper, we presented a robust a luminescence-based enzymatic assay applicable to an HTS setting (Fig. 1). When used to screen a large collection of 56,000 synthetic compounds, it only identified 9 initial hits and 5 of them were subsequently confirmed (Fig. 3 and Table 1). It appears that the hit rate against eEF2K in our screening system is rather low although the assay parameters may be explained by data from additional HTS campaigns with more diverse compound libraries.

Another interesting observation relates to inhibition of eEF2 phosphorylation by the newly discovered eEF2K inhibitors. Unlike the positive control (A484954), both WNN0048-A002 and WNN0017-C003 were unable to show such an effect, but instead, they promoted eEF2 phosphorylation in MDA-MB-453 breast cancer cells (Fig. 4). At this time we do not understand why this happened and whether it was due to a non-specific effect of these compounds or they hit other targets in the signaling pathways upstream of eEF2K. Extending the investigation to other confirmed hits may provide some clues. It should be noted that eEF2K is activated under a wide range of stress conditions.

To date, some small molecule eEF2K inhibitors have been reported. They are summarized and compared in terms of the experimental methods led to their discovery in Table 3. Rottlerin is mainly used as a selective inhibitor of protein kinase C_{δ}. It was found to activate calcium and potassium ion channels and to influence the function of mitochondria, thus can also be used as an eEF2K inhibitor. 1,3-selenazine analogs react with cysteine residues in cells thereby preventing eEF2 phosphorylation. NH125 is a widely accepted eEF2K inhibitor: it inhibits eEF2 activity in vitro but promotes eEF2 phosphorylation in vivo. Thieno [2,3-b] pyridine is an ATP-competitive inhibitor with weak inhibitory effects on eEF2K in cells. Thiopyran dicarboxonitrile analogs are a class of eEF2K inhibitors discovered by HTS. Finally, A484954 employed in this study is also an ATP-competitive inhibitor possessing a good inhibitory effect on eEF2K in vitro but rather low potency in cells. It is also cytotoxic at high concentrations.

Clearly, there exists an unmet demand for novel eEF2K inhibitors which can inhibit eEF2K effectively but have low or none cell cytotoxicity. In this regard, our work with eEF2K is just a beginning on the road to validate if it is an ideal target for cancer therapy.
<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Experimental method</th>
<th>IC₅₀ (µmol/L)</th>
<th>Ref.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Rottlerin</td>
<td><img src="image1.png" alt="Structure" /></td>
<td>Inhibitor of protein kinase Cα. <em>In vitro</em> eEF2K kinase assay followed by autoradiography. It inhibits various other protein kinases more strongly than eEF2K.</td>
<td>5.3</td>
<td>18</td>
</tr>
<tr>
<td>TS-2</td>
<td><img src="image2.png" alt="Structure" /></td>
<td>1,3-selenazine derivatives. <em>In vitro</em> eEF2K kinase assay followed by autoradiography.</td>
<td>0.36</td>
<td>19</td>
</tr>
<tr>
<td>TS-4</td>
<td><img src="image3.png" alt="Structure" /></td>
<td>Imidazolium histidine kinase inhibitors. <em>In vitro</em> eEF2K kinase assay followed by either a filter-based assay or immunoblotting.</td>
<td>0.06</td>
<td>20</td>
</tr>
<tr>
<td>NH125</td>
<td><img src="image4.png" alt="Structure" /></td>
<td>Thieno[2,3-b]pyridine analogues. Fluorometric, coupled-enzyme assay.</td>
<td>0.17</td>
<td>21</td>
</tr>
<tr>
<td>Compound 34</td>
<td><img src="image5.png" alt="Structure" /></td>
<td>Method 1: Fluorescence-based assay that uses the phosphorylation of a Sox-based peptide.</td>
<td>60</td>
<td>22</td>
</tr>
<tr>
<td>2,6-Diamino-4-(2-fluorophenyl)-4H-thiopyran-3,5-dicarbonitrile</td>
<td><img src="image6.png" alt="Structure" /></td>
<td>Method 2: Luminescence-based assay that measures the amount of ATP remaining in the reaction.</td>
<td>77</td>
<td>22</td>
</tr>
<tr>
<td>A-484954</td>
<td><img src="image7.png" alt="Structure" /></td>
<td>Discovered by Abbott Laboratories.</td>
<td>0.28</td>
<td>23</td>
</tr>
<tr>
<td>TX-1918</td>
<td><img src="image8.png" alt="Structure" /></td>
<td>2-Hydroxyaryliden-4-cyclopentene-1,3-diones analogues. <em>In vitro</em> eEF2K kinase assay followed by autoradiography. It also inhibits Src, PKA, PKC, and EGFR-K.</td>
<td>0.44</td>
<td>24</td>
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</table>
Acknowledgments

We are indebted to Claire Moore, Liang Qiu and Caihong Zhou for technical assistance. This work was partially supported by the National Health and Family Planning Commission of China (2012ZX09304-011, 2013ZX09401003-005, 2013ZX09507001 and 2013ZX09507-002), Shanghai Science and Technology Fund (15DZ2291600) and the Thousand Talents Program in China.

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