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User Guide

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Table of Contents

| | |
|---|---------------|
| 1. Pre-Analysis Instructions | - 3 - |
| 1.1 Dataset Requirements..... | - 3 - |
| 1.1.1 Z-Test/Kolmogorov-Smirnov | - 3 - |
| 1.1.2 KARP | - 4 - |
| 1.2 Analysis Parameters..... | - 4 - |
| 2. KSEAPIus Results..... | - 5 - |
| 2.1 Overview | - 5 - |
| 2.2 Kinase Activity Data..... | - 6 - |
| 2.2.1 Z-Test..... | - 6 - |
| 2.2.2 Kolmogorov-Smirnov | - 6 - |
| 2.2.3 KARP | - 7 - |
| 2.3 Kinase-Substrate Relationships | - 7 - |
| 2.4 Graphs | - 8 - |
| 2.4.1 Single-Sample Analysis..... | - 8 - |
| 2.2.1 Multiple-Sample Analysis..... | - 9 - |
| 3. Warnings | - 10 - |
| 4. Useful References..... | - 10 - |

1. Pre-Analysis Instructions

1.1 Dataset Requirements

- Please ensure only .tsv files no larger than 6MB are uploaded.
- Sample datasets are provided under the **Getting Started** tab of the KSEAPlus application.
- Each column must be titled and each row must have a value.
- The first column of a dataset must refer to the detected phosphosites, in the format shown below. Multiple sites may be listed on the same line but they must be separated by semicolons. The remaining columns must refer to the corresponding log₂(FC)/intensity values for each site. Different datasets are required depending on the algorithm chosen:

1.1.1 Z-Test/Kolmogorov-Smirnov

- These tests require columns 2 and onwards to contain pre-calculated log₂ transformed fold-change (FC) values for each site. Before taking the log₂ of the FC, it is generally advised to ensure the FC has been calculated in the format of *Experiment / Control* and not the opposite.

| | 1 | 2 | 3 | 4 |
|----|--------------------------|-------------------|-------------------|------------------|
| 1 | sh.index.sites | MCF7.GDC0941.fold | MCF7.AZD6482.fold | MCF7.CAL101.fold |
| 2 | KDM1A(T59); | 0.028609794 | 0.084441611 | -0.63413072 |
| 3 | INCENP(S143); | 0.400172957 | -0.103155816 | 0.645288689 |
| 4 | INCENP(S142); | 0.270414436 | -0.243416725 | 0.380882629 |
| 5 | INCENP(T135); | 0 | 0 | 0 |
| 6 | SAMD1(S161); | -1.172846259 | 0.151039761 | 0.937267317 |
| 7 | SAMD1(T157); | 0 | 0 | 0 |
| 8 | UNKL(S15); | -1.203364417 | -0.207652863 | 0.239831187 |
| 9 | NCL(S184);NCL(S206); | 0 | 0 | 0 |
| 10 | ARID3A(S77); | 2.238021203 | 0.445462684 | -0.274726094 |
| 11 | MSH6(S41); | -0.166792164 | 0.313911974 | -1.028279154 |
| 12 | PIPSL(S741);PSMD4(S256); | 6.091880369 | 2.270587187 | 6.560887118 |

1.1.2 KARP

- This test requires columns **2** and onwards to contain raw peptide intensity values for each identified phosphosite.

| | 1 | 2 | 3 | 4 |
|----|----------------------------|-----------------|-----------------|---------|
| 1 | gene; | AZD5438.3.PP2.1 | AZD5438.4.PP2.2 | DMSO.3 |
| 2 | KDM1A(T59); | 0.000216 | 0.000582 | 1.64017 |
| 3 | INCENP(S148); | 6.720432 | 7.867873 | 5.45268 |
| 4 | INCENP(T145); | 7.942432 | 9.098164 | 5.69186 |
| 5 | INCENP(S143); | 31.752216 | 37.470582 | 26.6093 |
| 6 | INCENP(S142); | 0.000324 | 0.000873 | 0.00051 |
| 7 | INCENP(M136); | 8.240108 | 21.65 | 14.4902 |
| 8 | INCENP(S143);INCENP(M136); | 1.52 | 2.31 | 0.8 |
| 9 | INCENP(T135); | 0.000216 | 0.000582 | 0.00034 |
| 10 | EIF3J(S11);EIF3J(S13); | 58.57 | 48.43 | 42.56 |
| 11 | RPL4(353-364, no_mod); | 0.000108 | 0.000291 | 0.00017 |
| 12 | SAMD1(S161); | 167.1 | 170.3 | 322.8 |

1.2 Analysis Parameters

- In order to upload a phosphoproteomic dataset, navigate to **Upload File** and select **Browse** to locate the .tsv file of interest.
- Click **Select kinase-substrate database** to choose the kinase-substrate relationships database for the dataset to be scanned against.
- Select a suitable KSEA algorithm from the **Select KSEA algorithm** dropdown list. Each algorithm can run against a single sample (e.g. one cell line) dataset or multiple samples (e.g. more than one cell line) dataset. Single sample analysis returns interactive and downloadable result tables and a barplot of scores for each kinase, and a multiple samples analysis returns a heatmap and downloadable .csv result files only.
- **Generate a barplot/heatmap?** allows to specify whether or not you would like to generate graphical kinase score results in the form of a barplot

(single sample analysis) or heatmap (multiple samples). Selecting **No** can speed up the analysis if the number of samples to analyse is large.

- **Plot parameters** - applicable to analyses with graphical output only (i.e. user has selected **Yes** to above). Minimum number of substrates determines which kinases to display on the barplot/heatmap. E.g. if 5 is selected, only kinases with 5 or more substrates will be displayed on the plots. KSEA results for the remaining kinases are calculated regardless, however they will not be shown on the plots.

2. KSEAPIus Results

2.1 Overview

The results section is divided into four main tabs:

- **Kinase Activity Data:** contains activity scores and related statistics for each identified kinase. This is presented as an interactive table for the single-sample analysis.
- **Kinase-Substrate Relationships:** contains additional information on identified unique kinase-substrate relationships, e.g. neighbouring residue sequences associated with a phosphosite, database source, and log₂(FC)/intensity value for each phosphosite. This is presented in the form of an interactive table for single sample analysis.
- **Graphs:** if graphical output is selected, this is where a plot will appear.
- **Download:** contains downloadable kinase activity data (.csv), K-S Relationships data (.csv) and barplot/heatmap (.svg).

2.2 Kinase Activity Data

Each KSEAPlus algorithm returns kinase activity scores and associated information for each kinase identified in the database search. The results may differ depending on the algorithm chosen, as outlined below:

2.2.1 Z-Test

- **Kinase:** gene corresponding to the identified kinase.
- **Sub.Count:** the number of substrates from the experimental dataset found for the kinase of interest.
- **mnlog2(FC):** mean $\log_2(\text{FC})$ across all of the kinase's substrates.
- **zSc:** Z-score denoting the activity level of the kinase. If the raw FC has been calculated in the format of Experiment / Control, a positive Z-score signifies an upregulated kinase activity in relation to the control. The opposite becomes true for negative Z-scores.
- **pVal:** p-value calculated based on the z-score. It denotes the statistical significance of the obtained result.

2.2.2 Kolmogorov-Smirnov (KS)

- **Kinase:** gene corresponding to the identified kinase.
- **Sub.Count:** the number of substrates from the experimental dataset found for the kinase of interest.
- **mnlog2(FC):** mean $\log_2(\text{FC})$ across all of the kinase's substrates.
- **(+/-) KS:** KS test statistic. It is signed based on the mean $\log_2(\text{FC})$ of the substrate group.
- **pVal:** p-value to denote the statistical significance.
- **(+/-) -log10(pVal):** $-\log_{10}$ of the p-value that is then signed based on the mean $\log_2(\text{FC})$ of the substrate group.

2.2.3 KARP

- **Kinase:** gene corresponding to the identified kinase.
- **Sub.Count:** the number of substrates from the experimental dataset found for the kinase of interest.
- **Total.Sub.Count:** the total number of substrates found for the kinase in the database of choice.
- **Sum.Ints:** the sum of all intensities across the substrate group.
- **kSc:** K-score denoting the activity level of the kinase.

2.3 Kinase-Substrate Relationships

- **Kinase:** gene associated with the identified kinase.
- **Site:** phosphosite phosphorylated by the corresponding kinase.
- ***Site.Seq (+/- 7AA):** A sequence of seven amino acids on either side of the affected phosphosite. Reported for KSEAPlus analyses against PhosphoSitePlus database only.
- **Source:** database source of the reported information.
- ***log2(FC):** log2(FC) value from the dataset associated with the identified phosphosite. Applicable only to Z-test and KS analyses.
- ***Ints.:** raw peptide intensity value from the dataset associated with the identified phosphosite. Applicable only to KARP analyses.

NOTE: if multiple samples analysis is selected, the column name specified by the user for each sample is appended to the relevant result columns in a repeating manner, e.g.:

| Kinase | Sub.Count | mnlog2(FC).CAL101 | zSc.CAL101 | pVal.CAL101 | mnlog2(FC).AZD6482 | zSc.AZD6482 | pVal.AZD6482 |
|--------|-----------|-------------------|--------------|-------------|--------------------|--------------|--------------|
| PLK1 | 1 | 0.645620068 | 0.118089438 | 0.452998396 | -0.155576994 | 0.017348215 | 0.493079411 |
| CDK1 | 4 | -0.179403447 | -0.074167351 | 0.470438612 | -0.449554994 | -0.085009667 | 0.466126853 |
| CDK2 | 2 | 0.221989856 | 0.054322327 | 0.478339181 | -0.216898798 | 0.006877713 | 0.497256211 |
| CDK5 | 1 | 2.376822021 | 0.443699547 | 0.328629911 | -2.625218176 | -0.485463405 | 0.313673833 |
| MTOR | 1 | 1.596297096 | 0.296895874 | 0.383273003 | 0.129651354 | 0.075419861 | 0.469940329 |
| DYRK1A | 1 | 0.208895522 | 0.035948861 | 0.485661568 | -0.261739751 | -0.004266207 | 0.498298035 |
| MAPK3 | 1 | -3.173241944 | -0.600174584 | 0.274194945 | -0.902877684 | -0.134799989 | 0.446385007 |

2.4 Graphs

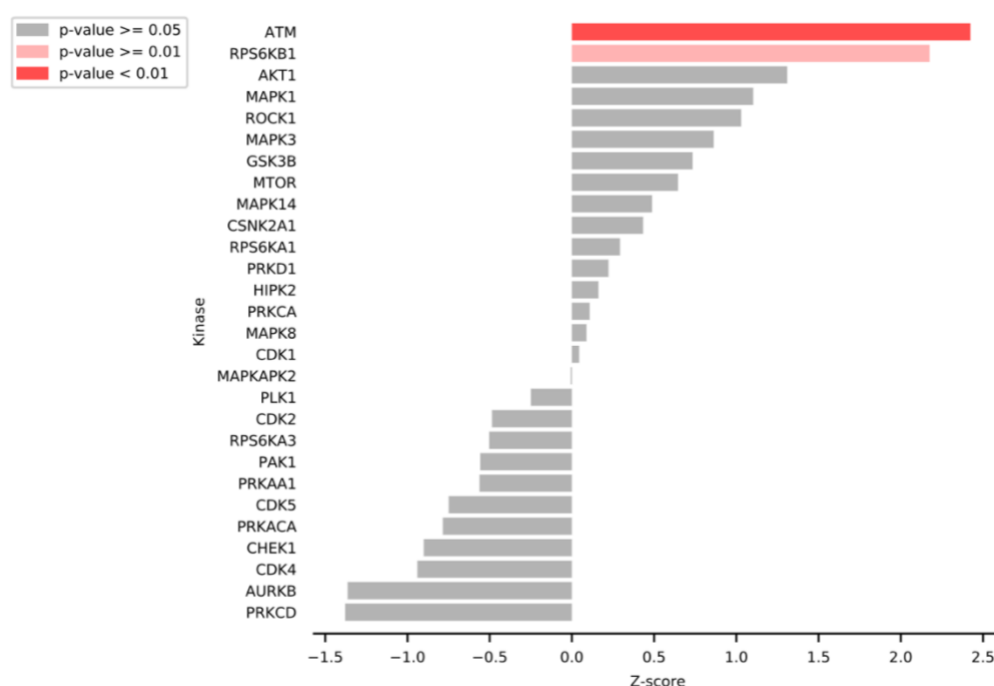
All kinase scores displayed on the plots are in the form of the Z-score, K-score and $-\log_{10}(\text{p-value})$ for the Z-test, KARP and KS test, respectively.

2.4.1 Single-Sample Analysis

Single-sample analysis returns a horizontal barplot containing scores corresponding to each identified kinase.

- For the Z-test and KS test, the bars are coloured differently depending on the statistical significance of the scores, i.e. bright red denotes p-values smaller than 0.01, pink indicates p-values greater than or equal to 0.01 and grey indicates p-values greater than or equal to 0.05.

An example barplot for a single-sample Z-test is shown below:



2.4.2 Multiple-Samples Analysis

Submitting a dataset containing multiple samples for analysis returns a heatmap of kinase scores corresponding to each kinase per each sample.

- **Asterisk annotation:** heatmap scores are further annotated for the Z-test and KS test if they are statistically significant (significance level 0.05). A single asterisk (*) denotes a p-value smaller than 0.05 and a double asterisk (**) indicates a p-value smaller than 0.01. No annotation is available for the KARP test.
- **Colour bar:** is shown on the right hand-side of the heatmap and indicates the magnitude of the scores. Red-like colours are associated with higher scores and blue-like colours with lower scores.

An example heatmap for a multiple samples Z-test analysis is shown below:

| | MCF7.AZD6482.fold | MCF7.CAL101.fold | MCF7.GDC0941.fold | MCF7.HS173.fold | MCF7.PIK294.fold | MCF7.TGX221.fold | MCF7.CX4945.fold | MCF7.Go6983.fold | MCF7.KN93.fold | MCF7.MK2206.fold |
|---------|-------------------|------------------|-------------------|-----------------|------------------|------------------|------------------|------------------|----------------|------------------|
| PLK1 | -0.2 | 0.7 | -0.1 | 0.4 | -0.2 | 0.9 | 1.7* | 1.3 | 0.8 | 2.0* |
| CDK1 | 0.0 | 1.8* | 1.2 | 2.1* | 1.3 | 1.3 | -2.3* | -1.1 | -1.7* | -2.4** |
| CDK2 | -0.5 | 0.9 | 0.6 | 0.2 | 0.8 | 0.5 | -0.5 | -0.2 | -0.4 | -1.1 |
| CDK5 | -0.7 | 0.4 | 0.9 | 0.1 | 0.2 | 0.3 | -1.6 | -1.4 | -0.6 | -1.7* |
| MTOR | 0.6 | 1.0 | -2.5** | -0.8 | 0.9 | 0.9 | 0.5 | 0.4 | 0.2 | -1.4 |
| MAPK3 | 0.9 | 0.9 | 0.3 | 1.5 | 0.9 | 1.1 | -1.7* | -1.1 | -0.3 | 0.1 |
| CSNK2A1 | 0.4 | -0.9 | 1.2 | 1.2 | -0.5 | 0.2 | 5.7** | 4.6** | 4.8** | 6.2** |
| MAPK1 | 1.1 | 1.0 | 0.2 | 1.4 | 1.1 | 1.0 | -1.3 | -1.2 | -0.6 | -0.9 |

3. Warnings

- **Request timeout:** KSEAPlus is running on a free Heroku dyno where a request automatically times out after 30 seconds if no data have been sent to the browser. When uploading a relatively large dataset (e.g. ~5MB) with multiple columns, consider omitting the heatmap generation as this can significantly slow down the analysis.
- **File size limit:** current version of KSEAPlus accepts .tsv files up to 6 MB in size.
- **Downloads/session timeout:** for each session, all of the generated results data is stored temporarily in an in-memory server-side key-value store (Heroku-Redis). To preserve space, every session expires after 10 minutes and its data is deleted therefore please ensure you download your results before the session expires.

4. Useful References

Hornbeck, P., Zhang, B., Murray, B., Kornhauser, J., Latham V., Skrzypek E. PhosphoSitePlus, 2014: mutations, PTMs and recalibrations. *Nucleic Acids Res.* 2015 43:D512-20.

Wiredja, D., Koyutürk, M. and Chance, M. (2017). The KSEA App: a web-based tool for kinase activity inference from quantitative phosphoproteomics. *Bioinformatics*, 33(21), pp.3489-3491.

Casado, P., Rodriguez-Prados, J., Cosulich, S., Guichard, S., Vanhaesebroeck, B., Joel, S. and Cutillas, P. (2013). Kinase-Substrate Enrichment Analysis Provides Insights into the Heterogeneity of Signaling Pathway Activation in Leukemia Cells. *Science Signaling*, 6(268).

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