Supplementary manuscript of

Identifying personalized driver genes of cancer individual patients with semi-supervised graph neural network

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## Supplementary note 1: The statistic information of positive and negative genes in PGIN

Table S1 The statistic information of positive and negative genes in PGIN

|  |  |  |  |
| --- | --- | --- | --- |
|  | BRCA | LUSC | LUAD |
| #posi | 18.4602+-1.5749 | 27.7292+-3.0479 | 29.4286+-3.7005 |
| #neg | 191.9115+-15.8389 | 419.7292+-23.0389 | 412.5000+-24.4861 |

## Supplementary note 2: details of class weighted loss function

To balance the loss of imbalanced classes or labels in PGIN, we adopted a class-balanced softmax cross-entropy loss function by introducing a normalized weighting factor,1/where is the effective number of genesfor class *y*in PGIN. The effective number of genes for class *y* is defined as ),=*N*/(*N*-1) where *ny* is the number of genes in the class or label *y* and *N* is the number of genes in PGIN. The class weighted loss function can be written as



wheredenotes the model's estimated class probabilities. The weighting factor is a normalized vector=*C; C* is the number of classes (here *C* = 2); is softmax cross-entropy loss function. If the gene *x*is within a positive label set, its label , otherwise, .

## Supplementary note 3: Experimental parameters sensitivity analysis

To further demonstrate the effectiveness of the setting parameters in our PersonalizedGNN, we tuned one of the parameters over suitable ranges of parameter values while we fixed other parameters in BRCA, LUSC, LUAD data sets. For each cancer data set, we used the same parameters among all individual patients andcalculated the average gene ranking score among all individual patients for each cancer data. Based on the average gene ranking score of the cancer data, we obtained the gene ranking result and calculated the *AveragePrecision* of different parameter choices on each cancer data. The parameter setting resultson BRCA, LUAD, and LUSC were shown in **Fig S1 of Additional file 1**. As shown in Figure S1, PersonalizedGNN has the best performance for our setting parameters in PersonalizedGNN.

**para check**

**Fig S1**The error bar in terms of *AveragePrecision*for different parameter choices among top *k*(*k*=1,2,...,30) ranking genes on BRCA, LUSC, LUAD data sets.

## Supplementary note 4: parameter setting of PersonalizedGNN and other methods

* **The parameters in PersonalizedGNN**

Table S2 The parameters of PersonalizedGNN on cancer driver gene prediction

|  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- |
| The number of layers | The number of nodes in each layer | weight decay | learn rate | coefficient of Dropout | coefficient of DropEdge | coefficient of DropAttention | The number of attention heads | epoch |
| 3 | 750 | 1E-7 | 0.02 | 0.25 | 0.75 | 0.75 | 5 | 2000 |

* **Parameters in other methods**

ChebNet: the graph convolution kernel size is 2; the number of layers is 3;the size of each hidden layer is 750; the number of epochs is 2000; bias in graph convolution is enabled, the activation function is ReLU; the dropout rate is 0.25; the learning rate is 0.02, and the optimizer is Adam.

GraphSAGE: the number of layers is 3; the size of each hidden layer is 750; the number of epochs is 2000; the number of sampled neighbors is 10; the batch size is 50; and the learning rate is 0.02.

GCN: the number of layers is 3; the size of hidden layers is 750; the num of epochs is 2000; the the dropout rate is 0.25; learning rate is 0.02; and the weight decay is 1e-7.

GAT without label reuse: the heads of attention mechanism is 5; the number of layers is 3; the size of each hidden layer is 500; the number of epochs is 2000; the coefficient of Dropout, DropEdge and DropAttention is 0.25, 0.75 and 0.75 respectively; the learning rate is 0.02; the weight decay is 1e-7; and the activation function is LeakyReLU with the slope alpha equals 0.2.

## Supplementary note 5: supplementary results of PersonalizedGNN

342b84516eba339c4ea137fec87cc6a4

**Fig.S2** Enrichment pathway results of NPDGson different subtypes of BRCA cancer patients.

770023a6d07613f2e38a5fa451d25461

**Fig.S3** Enrichment pathway results of NPDGson different subtypes of LUNG cancer patients.

## Supplementary note 6: details of in vitro cell-based assays

**Cell lines and cell culture**

LUSC cell line H1703 was purchased from Procell Life Science & Technology (Wuhan, China) and cultured in RPMI 1640 medium (GIBCO, USA) supplemented with 10% fetal bovine serum (FBS, BI, USA), 100 U/mL penicillin (Solarbio, China), and 100 μg/mL streptomycin (Solarbio, China). All cells were cultured at 37◦C in a humidified atmosphere of 95% air and 5% CO2.

**Plasmid construction and lentiviral transfection**

Short hairpin RNA (shRNA) was cloned into PLKO.1 vector and transfected into H1703(H1703 shFZD7) cells using a lentiviral transfection system. The empty vector was transfected into H1703 cells as a control (H1703 shNT). The target sequence is 5’-GCCGCTTCTACCACAGACTTA-3'.

**qRT‑PCR analysis**

According to the manual, total RNA was extracted from cell lines and tissue samples using a total RNA isolation kit (Tiangen, DP419, China). Tefrst strand cDNA was synthesized from 2 µg of total RNA using the Reverse Assisted First Strand cDNA Synthesis Kit (ThermoFisher, K1622, USA). The qRT-PCR assay was performed using SYBR Green I Master (Roche, 04887352001, Switzerland) in Roche LightCycler 480 II. The primers for FZD7 were 5’-CGTGTCGTTCTCTGTGCGAG-3'and 5'-GTAGGGCGCGGTAGGGTAG-3'. *GAPDH* was used as a control to normalize the expression of each gene.

**Cell viability assay**

H1703 cells (3000 cells/well) were seeded in 96-well plates and cultured overnight. MTT (Sigma) was added to the culture wells at 24 h, 48 h, 72 h, 96 h, and 120 h and incubated at 37 °C for 4 h. After incubation, the medium was discarded, and 150 μL of dimethyl sulfoxide was added to each well to dissolve the formazan crystals. The absorbance at 490 nm was measured using a microplate reader.

**Colony formation assay**

The siRNA-transfected H1703 cells were seeded in six-well plates with 3000 cells/well and cultured for 7 days. Then the medium was aspirated, and the cells were washed with phosphate-buffered saline (PBS) and fixed with 4% paraformaldehyde solution at room temperature for 30 min. Thereafter, the fixative was removed, and 0.2% crystal violet staining solution was added to stain the cells at room temperature for 30 min. After staining, the crystal violet was rinsed with PBS, and the cells were dried at room temperature. Then, the colonies of cells were counted and photographed.

**Invasion assay**

In vitro invasion assays were performed using Transwell chambers (Corning, MA, USA). H1703 cells (1×105) transfected with shRNA and negative shRNA were plated on the 24-well plate and cultured for 24 h. After the incubation, the medium was discarded and fixed at room temperature for 30 min. Then, 0.2% crystal violet staining solution was added to stain the cells at room temperature for 30 min. After staining, the chamber was washed with PBS, and the cells were carefully wiped off with a cotton swab. The stained cells were counted under a microscope and photographed.

**Western blot assay**

Anti-FZD7 antibody (A4213), anti-CyclinA2antibody (A7632), and anti-CyclinE2 antibody (A9305) were purchased from ABclonal (Cambridge, MA, USA). Anti-P27 antibody (25614-1-AP) was purchased from Abcam. Anti-β-actin (GB11001) antibody was purchased from Servicebio. Total cellular protein was extracted by adding a protein lysis buffer. Proteins were separated by sodium dodecyl sulfate-polyacrylamide gel electrophoresis and transferred to a polyvinylidene difluoride (PVDF) membrane (Merck Millipore, IPVH00010, USA). The membrane was blocked with 5% milk dissolved in Tris-buffered saline (TBST) for 1 h at room temperature. Then, the PVDF membrane was incubated with the antibody (1:1000)dissolved in 5% milk overnight at 4 °C. After that, the membrane was washed six times for 5 min each time using TBST. Thereafter, the membrane was incubated with horseradish peroxidase-conjugated mouse anti-rabbit antibody (1:8,000) for 2 h at room temperature. Finally, the membrane was washed six times, and the enhanced chemiluminescence detection kit (Applygen Technologies, Beijing, China) was used to expose the protein bands. The results were analyzed using ImageJ software (NIH, Bethesda, MD, USA).

**Cell cycle assays**

The cell cycle was analyzed using a Cell Cycle Staining Kit (eBioscience). Each sample with 5 × 106 cells was washed with PBS and incubated with 500 μL DNA staining solution and 5 μL permeabilization solution for 30 min at room temperature. The samples were analyzed by flow cytometry.