

TFE3 Rabbit mAb

Catalog No.: A29447 **Recombinant**

Basic Information

Observed MW

72-89 kDa

Calculated MW

11 kDa/62 kDa

Category

Primary antibody

Applications

WB, IP, IHC-P, ChIP, ELISA

Cross-Reactivity

Human, Mouse, Rat

CloneNo number

ARC4257

Background

This gene encodes a basic helix-loop-helix domain-containing transcription factor that binds MUE3-type E-box sequences in the promoter of genes. The encoded protein promotes the expression of genes downstream of transforming growth factor beta (TGF-beta) signaling. This gene may be involved in chromosomal translocations in renal cell carcinomas and other cancers, resulting in the production of fusion proteins. Translocation partners include PRCC (papillary renal cell carcinoma), NONO (non-POU domain containing, octamer-binding), and ASPSCR1 (alveolar soft part sarcoma chromosome region, candidate 1), among other genes. Alternative splicing results in multiple transcript variants.

Recommended Dilutions

WB 1:3000 - 1:15000**IP** 0.5 µg - 4 µg antibody for
200 µg - 400 µg extracts of
whole cells**IHC-P** 1:500 - 1:2000**ChIP** 2 µg antibody for 15 µg - 20
µg of Chromatin**ELISA** Recommended starting
concentration is 1 µg/mL.
Please optimize the
concentration based on your
specific assay requirements.
For high-ratio antibody
dilutions ($\geq 1:10000$) a
sequential dilution method
is strongly recommended to
ensure measurement
accuracy.

Immunogen Information

Gene ID

7030

Swiss Prot

P19532

Immunogen

This information is considered to be commercially sensitive.

Synonyms

TFEA; RCCP2; RCCX1; MRXSPF; bHLHe33

Product Information

Source

Rabbit

Isotype

IgG

Purification

Affinity purification

Storage

Store at -20°C. Avoid freeze / thaw cycles.

Buffer: PBS with 0.09% Sodium azide, 0.05% BSA, 50% glycerol, pH7.3.

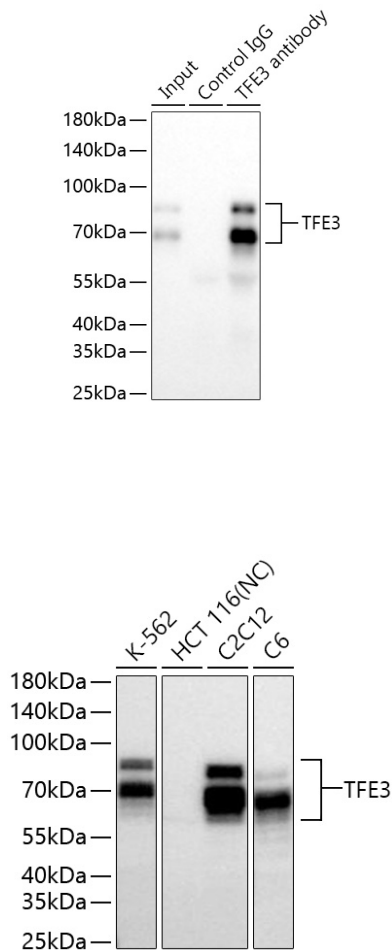
Contact

☎ | 400-999-6126

✉ | cn.market@abclonal.com.cn

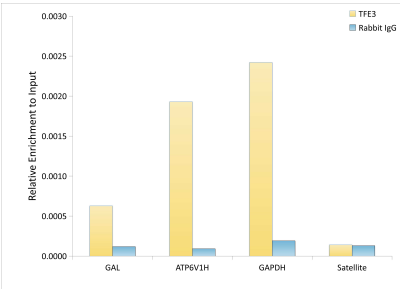
🌐 | www.abclonal.com.cn

Validation Data

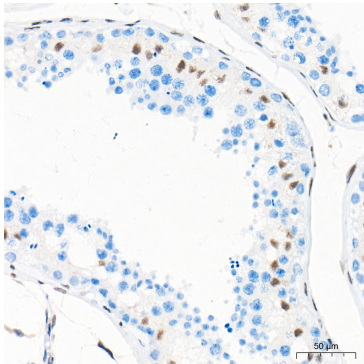
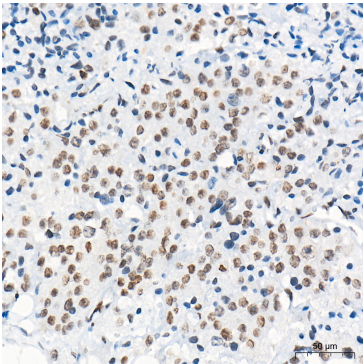
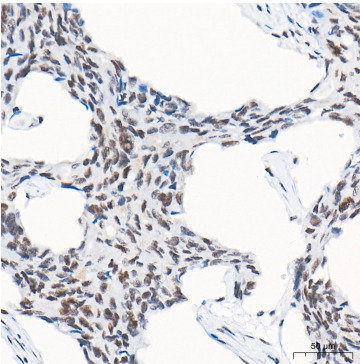


Immunoprecipitation of TFE3 from 300 µg extracts of K-562 cells was performed using 2 µg of TFE3 Rabbit mAb (A29447). Rabbit Control IgG (AC005) was used to precipitate the Control IgG sample. IP samples were eluted with 1x Laemmli Buffer. The Input lane represents 10% of the total input. Western blot analysis of immunoprecipitates was conducted using TFE3 Rabbit mAb (A29447) at a dilution of 1:5000.

Western blot analysis of various lysates using TFE3 Rabbit mAb (A29447) at 1:5000 dilution incubated overnight at 4°C. Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution. Lysates/proteins: 25 µg per lane. Blocking buffer: 3% nonfat dry milk in TBST. Detection: ECL Basic Kit (RM00020). Negative control (NC): HCT 116. Exposure time: 10 s.

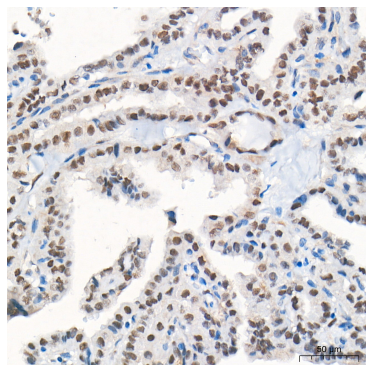


Chromatin immunoprecipitation was performed with 20 µg of cross-linked chromatin from K-562 cells, using 2 µg of TFE3 Rabbit mAb (A29447) and Rabbit IgG isotype control (AC042). The enrichment of immunoprecipitated DNA at different genomic loci was examined by quantitative PCR. The histogram compares the ratio of the immunoprecipitated DNA to the input at given loci.



Validation Data

Immunohistochemistry analysis of paraffin-embedded Human breast cancer tissue using TFE3 Rabbit mAb (A29447) at a dilution of 1:700 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.



Immunohistochemistry analysis of paraffin-embedded Human thyroid cancer tissue using TFE3 Rabbit mAb (A29447) at a dilution of 1:700 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.

Immunohistochemistry analysis of paraffin-embedded Human prostate cancer tissue using TFE3 Rabbit mAb (A29447) at a dilution of 1:700 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.

Immunohistochemistry analysis of paraffin-embedded Human testis tissue using TFE3 Rabbit mAb (A29447) at a dilution of 1:700 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.