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NKX3.1 (D6D2Z) XP® Rabbit mAb



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For Research Use Only. Not for Use in Diagnostic Procedures.

Applications:Reactivity:Sensitivity:MW (kDa):Source/Isotype:UniProt ID:Entrez-Gene Id:WB, IHC-PHEndogenous30Rabbit IgG#Q998014824

Product Usage
InformationApplicationDilution
1:1000Western Blotting
Immunohistochemistry (Paraffin)1:250

 $\textbf{Storage} \hspace{1.5cm} \textbf{Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 } \mu\text{g/ml BSA, 50\% glycerol and less than} \\$

0.02% sodium azide. Store at -20° C. Do not aliquot the antibody.

For a carrier free (BSA and azide free) version of this product see product #32253.

Specificity / Sensitivity

NKX3.1 (D6D2Z) XP® Rabbit mAb recognizes endogenous levels of total NKX3.1 protein.

Source / Purification

Monoclonal antibody is produced by immunizing animals with recombinant protein specific to the amino terminus of human NKX3.1 protein. The epitope is near the amino terminus, in a region that is 100% conserved between isoforms 1 and 5 of human NKX3.1.

Background

NKX3.1 is a homeobox transcription factor that in mammals plays a defining role in embryonic prostate morphogenesis. The expression of mammalian NKX3.1 is androgen-dependent, restricted primarily to developing and mature prostate epithelium, and is frequently reduced or lost in prostate cancer (1-3). The human *NKX3.1* gene is located on chromsome 8p21.2, within a region that shows loss of heterozygosity (LOH) in >50% of prostate cancer cases (2). Allelic loss at the *NKX3.1* locus is also common in high grade Prostate Intraepithelial Neoplasia (PIN), thought to be a putative precursor lesion to invasive prostate adenocarcinomas, suggesting that LOH at the *NKX3.1* locus is a critical early step in prostate cancer development (4). Notably, the remaining *NKX3.1* allele is intact in the majority of LOH cases, leading to the suggestion that NKX3.1 functions as a haploinsufficient tumor suppressor (4-6). Due to its highly restricted expression in prostate epithelial cells, NKX3.1 has been suggested as a diagnostic marker of prostate carcinoma (7), and may have additional utility as a biomarker of metastatic lesions originating in the prostate (8).

Background References

- 1. Bhatia-Gaur, R. et al. (1999) Genes Dev 13, 966-77.
- 2. He, W.W. et al. (1997) Genomics 43, 69-77.
- 3. Bowen, C. et al. (2000) Cancer Res 60, 6111-5.
- 4. Magee, J.A. et al. (2003) Cancer Cell 3, 273-83.
- 5. Voeller, H.J. et al. (1997) *Cancer Res* 57, 4455-9.
- 6. Bethel, C.R. et al. (2006) Cancer Res 66, 10683-90.
- 7. Epstein, J.I. et al. (2014) Am J Surg Pathol 38, e6-e19.
- 8. Conner, J.R. and Hornick, J.L. (2015) Adv Anat Pathol 22, 149-67.

Species Reactivity

Species reactivity is determined by testing in at least one approved application (e.g., western blot).

Western Blot Buffer

 $IMPORTANT: For western \ blots, incubate \ membrane \ with \ diluted \ primary \ antibody \ in \ 5\% \ w/v \ nonfat \ dry$

milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

Applications Key

WB: Western Blotting IHC-P: Immunohistochemistry (Paraffin)

Cross-Reactivity Key

H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus Mi: mink C: chicken Dm: D. melanogaster

X: Xenopus Z: zebrafish B: bovine Dg: dog Pg: pig Sc: S. cerevisiae Ce: C. elegans Hr: horse

GP: Guinea Pig Rab: rabbit All: all species expected

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