



## Human FOXN1 peptide (DAG-P0526)

This product is for research use only and is not intended for diagnostic use.

### PRODUCT INFORMATION

<b>Antigen Description</b>	Mutations in the winged-helix transcription factor gene at the nude locus in mice and rats produce the pleiotropic phenotype of hairlessness and athymia, resulting in a severely compromised immune system. This gene is orthologous to the mouse and rat genes and encodes a similar DNA-binding transcription factor that is thought to regulate keratin gene expression. A mutation in this gene has been correlated with T-cell immunodeficiency, the skin disorder congenital alopecia, and nail dystrophy. Alternative splicing in the 5' UTR of this gene has been observed. [provided by RefSeq, Jul 2008]
<b>Specificity</b>	Expressed in thymus.
<b>Conjugate</b>	Unconjugated
<b>Sequence Similarities</b>	Contains 1 fork-head DNA-binding domain.
<b>Format</b>	Liquid
<b>Preservative</b>	None
<b>Storage</b>	Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.

### GENE INFORMATION

<b>Gene Name</b>	<a href="#">FOXN1 forkhead box N1 [ Homo sapiens (human) ]</a>
<b>Official Symbol</b>	FOXN1
<b>Synonyms</b>	FOXN1; forkhead box N1; WHN; RONU; FKHL20; forkhead box protein N1; Rowett nude; winged helix nude; winged-helix nude; winged-helix transcription factor nude;
<b>Entrez Gene ID</b>	<a href="#">8456</a>

<b>mRNA Refseq</b>	<a href="#">NM_003593.2</a>
<b>Protein Refseq</b>	<a href="#">NP_003584.2</a>
<b>UniProt ID</b>	O15353
<b>Chromosome Location</b>	17q11-q12
<b>Function</b>	DNA binding, bending; RNA polymerase II distal enhancer sequence-specific DNA binding transcription factor activity; double-stranded DNA binding; sequence-specific DNA binding; transcription factor binding;