

Artemis (pS516) Antibody

Purified Rabbit Polyclonal Antibody (Pab)
Catalog # AP51158

Product Information

Application	WB
Primary Accession	Q96SD1
Reactivity	Human, Mouse, Rat
Host	Rabbit
Clonality	Polyclonal
Calculated MW	78436

Additional Information

Gene ID	64421
Other Names	Protein artemis, 31--, DNA cross-link repair 1C protein, Protein A-SCID, SNM1 homolog C, hSNM1C, SNM1-like protein, DCLRE1C, ARTEMIS, ASCID, SCIDA, SNM1C
Target/Specificity	KLH-conjugated synthetic peptide encompassing a sequence within the center region of human Artemis. The exact sequence is proprietary.
Dilution	WB~ 1:1000
Format	0.01M PBS, pH 7.2, 0.09% (W/V) Sodium azide, Glycerol 50%
Storage	Store at -20 °C. Stable for 12 months from date of receipt

Protein Information

Name	DCLRE1C (HGNC:17642)
Function	Nuclease involved in DNA non-homologous end joining (NHEJ); required for double-strand break repair and V(D)J recombination (PubMed: 11336668 , PubMed: 11955432 , PubMed: 12055248 , PubMed: 14744996 , PubMed: 15071507 , PubMed: 15574326 , PubMed: 15936993). Required for V(D)J recombination, the process by which exons encoding the antigen-binding domains of immunoglobulins and T-cell receptor proteins are assembled from individual V, (D), and J gene segments (PubMed: 11336668 , PubMed: 11955432 , PubMed: 14744996). V(D)J recombination is initiated by the lymphoid specific RAG endonuclease complex, which generates site specific DNA double strand breaks (DSBs) (PubMed: 11336668 , PubMed: 11955432 , PubMed: 14744996). These DSBs present two types of DNA end structures: hairpin sealed coding ends and phosphorylated blunt signal ends (PubMed: 11336668 , PubMed: 11955432 , PubMed: 14744996). These ends are independently repaired by the non homologous end joining (NHEJ)

pathway to form coding and signal joints respectively (PubMed:[11336668](#), PubMed:[11955432](#), PubMed:[14744996](#)). This protein exhibits single-strand specific 5'-3' exonuclease activity in isolation and acquires endonucleolytic activity on 5' and 3' hairpins and overhangs when in a complex with PRKDC (PubMed:[11955432](#), PubMed:[15071507](#), PubMed:[15574326](#), PubMed:[15936993](#)). The latter activity is required specifically for the resolution of closed hairpins prior to the formation of the coding joint (PubMed:[11955432](#)). Also required for the repair of complex DSBs induced by ionizing radiation, which require substantial end-processing prior to religation by NHEJ (PubMed:[15456891](#), PubMed:[15468306](#), PubMed:[15574327](#), PubMed:[15811628](#)).

Cellular Location	Nucleus
Tissue Location	Ubiquitously expressed, with highest levels in the kidney, lung, pancreas and placenta (at the mRNA level). Expression is not increased in thymus or bone marrow, sites of V(D)J recombination

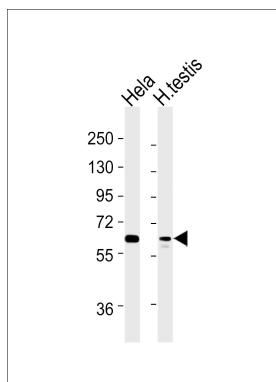
Background

Required for V(D)J recombination, the process by which exons encoding the antigen-binding domains of immunoglobulins and T-cell receptor proteins are assembled from individual V, (D), and J gene segments. V(D)J recombination is initiated by the lymphoid specific RAG endonuclease complex, which generates site specific DNA double strand breaks (DSBs). These DSBs present two types of DNA end structures: hairpin sealed coding ends and phosphorylated blunt signal ends. These ends are independently repaired by the non homologous end joining (NHEJ) pathway to form coding and signal joints respectively. This protein exhibits single-strand specific 5'-3' exonuclease activity in isolation and acquires endonucleolytic activity on 5' and 3' hairpins and overhangs when in a complex with PRKDC. The latter activity is required specifically for the resolution of closed hairpins prior to the formation of the coding joint. May also be required for the repair of complex DSBs induced by ionizing radiation, which require substantial end-processing prior to religation by NHEJ.

References

Moshous D.,et al.Cell 105:177-186(2001).
 Li L.,et al.J. Immunol. 168:6323-6329(2002).
 Ota T.,et al.Nat. Genet. 36:40-45(2004).
 Deloukas P.,et al.Nature 429:375-381(2004).
 Mural R.J.,et al.Submitted (SEP-2005) to the EMBL/GenBank/DDBJ databases.

Images



All lanes : Anti-Artemis (pS516) Antibody at 1:1000 dilution
 Lane 1: Hela whole cell lysates Lane 2: H.testis whole cell lysates Lysates/proteins at 20 µg per lane.
 Secondary Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/10000 dilution Predicted band size : 78 kDa Blocking/Dilution buffer: 5% NFDM/TBST.

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.