

### Anti-Calnexin antibody, rabbit polyclonal

73-026 100 μl

**Key words:** Canx, Endoplasmic reticulum, Chaperon, Synaptic vesicle endocytosis, Transmembrane, Calcium ion binding

**Function:** Calcium-binding protein that interacts with newly synthesized glycoproteins in the endoplasmic reticulum. It may act in assisting protein assembly and/or in the retention within the ER of unassembled protein subunits. It seems to play a major role in the quality control apparatus of the ER by the retention of incorrectly folded proteins. Associated with partial T-cell antigen receptor complexes that escape the ER of immature thymocytes, it may function as a signaling complex regulating thymocyte maturation. Additionally it may play a role in receptor-mediated endocytosis at the synapse

**Molecular mass:** 67,278 Da with 591 amino acids. It undergoes palmitoylation and phosphorylation.

#### Applications:

- 1. Western blotting (1/1,000 dilution))
- 2. Immunoprecipitation (1/100 dilution).
- 3. Immunohistochemistry (1/100 1/1,000 dilution)
- 4. Immuno-fluorescent staining (1/100~1/300 dilution)

**Immunogen:** C-terminal peptide of mouse Calnexin protein, EDEILNRSPRNRKPRRE, conjugated with KLH

Reactivity: Mouse, rat, human

Form: Whole rabbit antiserum added with 0.1% sodium azide.

**Storage:** Shipped at 4°C or at -20°C. Upon arrival, spin-down and store at -20°C for longer period.

Database Links: <u>uniprot/P35564</u> mouse calnexin. <u>Gene ID 12330</u> mouse calnexin.

Reference: This antibody was described in Ref.1 and used in the following publications. Ikawa M. et al.

(2001) Calmegin Is Required for Fertilin  $\alpha/\beta$  Heterodimerization and Sperm Fertility. <u>Dev Biol.</u> 240: 254-61.

**IP.** Open access.

 Ikawa M. et al. (2011) Calsperin is a testis-specific chaperone required for sperm fertility. <u>J Biol</u> <u>Chem.</u> 286: 5639-46. WB, IP. Open access.



Fig.1 Western blotting analysis of mouse testis extracts of different ages with anti-Calnexin (CANX) antibody.

20  $\mu$  g of Triton X-100 extracts from mouse testis was reacted with anti-Calnexin antiserum at 1/1,000 dilution. Calnexin expression started at birth. (wks) stands for weeks.



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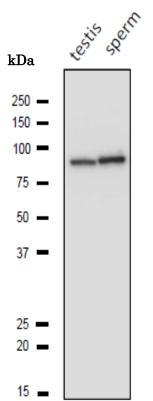


Fig.2 Western blotting analysis of lysatess of mouse testis and sperm with anti-Calnexin (CANX) antibody.

Proteins of the lysates (10 µg) of mouse testis and sperm were separated on SDS-PAGE and blotted to PVDF membrane and reacted with anti-Calnexin antiserum at 1/1,000 dilution. Calnexin bands migrated to the position corresponding to molecular mass of 90 kDa.

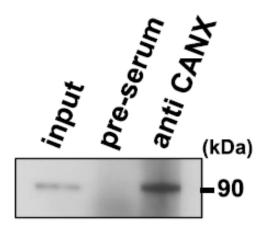
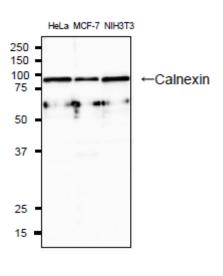


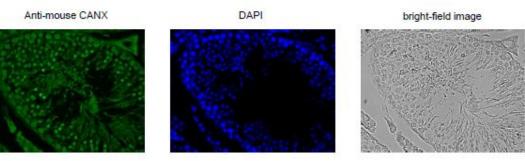
Fig.3.Immunoprecipitation of Calnexin from mouse testis. One mg of testis lysate (supernatant in lysis buffer containing 10 m m Tris-HCl pH8.0, 50 mM NaCl, 1% protease inhibitor mix) was incubated with 2  $\mu$ l of anti-Calnexin (CANX) antiserum and 50  $\mu$ l. of protein-A conjugated magnetic beads (Miltenyi Biotec) and immunoprecipitated according to the protocol of supplier. The immunoprecipitated sample was analyzed by western blotting using the same antibody at 1/1,000 dilution.





# Fig.4 Western blotting analysis of lysatess of human and mouse cell lines with anti-Calnexin (CANX) antibody.

Proteins of the lysates (20 µg) of HeLa, MCF7 and NIH3T3 cells were separated on SDS-PAGE and blotted to PVDF membrane and reacted with anti-Calnexin antiserum 1/1,000atdilution. As second antibody, goat polyclonal antibody to rabbit IgG conjugated with HRP (ab97051) was used



### Fig.5. Immunohistochemistry of mouse testis using anti-CANX antibody.

Formalin-fixed and paraffin-embedded mouse testis Deparaffinization by LemosolRA (#122-03991,Wako, Osaka)

Rehydration	100% Et-OH, 95%, 90%, 70%, DW				
Antigen retrieva	al Histo/Zyme (Cat.# k046; Diagnostic BioSystems)				
Washing	PBST (0.25% triton X-100/PBS-)				
Blocking	10 % FBS / PBST 30 min				
1st antibody 100 dilution in PBS- 4°C O/N					
Washing	PBS-				
2nd antibody	1000 dilution, 60 min (Alexa Flour-488 goat anti-rabbit IgG (H&L),				
	#1166843 ; molecular probes)				
washing PBS-	5 min X 3				
DAPI 1.0 µ g/mL DAPI in TBS 10 min					
(×100 stock, 0.1mg/mL)					
Washing	PBS-				
Mount ImmunoSelect Antifading Mounting Medium (SCR-38447; Dianova)					



## Fig.6 Immunofluorescent staining of HeLa cells with anti-Calnexin antibody

	Anti-mous	se CANX		DAPI	Merge		
1/100 dilution		4		•			
1/300 dilution				• • •			
	wash	PBS-, 2time		10° over night			
	fixed wash	4 % PFA 1. PBS- 5 min		/4℃ over night			
	permeabalized 0.25 % Triton X-100/PBS 10 min						
	wash PBS- , 5 min X 3 times						
	blocking 1.5% BSA/PBS, 30 min						
	1st antibody dilution buffer; 1.5 % BSA/PBS or CanGet signal B 100, 300 or 1000 dilution, 4℃ over night (r.t. for 60 min)						
	wash	PBS-, 5 min X 3 times					
	2nd antibody	1000 dilution, 60 min (Alex Fluor 488 goat-anti-rabbit IgG (H&L); molecular probes)					
	wash	PBS-, 5 min X 3 times					
	DAPI	1.0µg/mL DAPI in TBS, 5 min (X100 stock, 0.1mg/mL)					
	wash	0.1 % Triton X-100/TBS, 5min					
	TBS, 5min						
	mount ImmunoSelect Antifading Mounting Medium (SCR-38447; Dianova)						