

FAK (phospho Ser910) antibody

Cat. No. GTX24794

Host	Rabbit
Clonality	Polyclonal
Isotype	IgG
Applications	WB, FCM, IP
Reactivity	Human, Mouse

Package 50 μl

Applications

Application Note

*Optimal dilutions/concentrations should be determined by the researcher.

Suggested dilution	Recommended dilution
WB	Assay dependent
FCM	1:20
IP	Assay dependent
Not tosted in other applications	

Not tested in other applications.

Calculated MW 119 kDa. (Note)

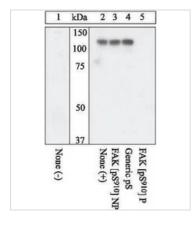
Properties	
Form	Liquid
Buffer	PBS, 0.1% BSA, 50% Glycerol
Preservative	0.05% Sodium azide
Storage	Store as concentrated solution. Centrifuge briefly prior to opening vial. For short-term storage (1-2 weeks), store at 4°C. For long-term storage, aliquot and store at -20°C or below. Avoid multiple freeze-thaw cycles.
Concentration	Batch dependent (Please refer to the vial label for the specific concentration.)
Immunogen	The antiserum was produced against a chemically synthesized phosphopeptide derived from the region of human FAK that contains serine 910. The sequence is conserved among multiple species including mouse, rat, chicken and frog.
Purification	Purified by antigen-affinity chromatography
Conjugation	Unconjugated
Note	For laboratory research use only. Not for any clinical, therapeutic, or diagnostic use in humans or animals. Not for animal or human consumption.
	Purchasers shall not, and agree not to enable third parties to, analyze, copy, reverse engineer or otherwise attempt to determine the structure or sequence of the product.



For full product information, images and publications, please visit our <u>website</u>.

Date 2025 / 12 / 27 Page 1 of 2

DATA IMAGES



GTX24794 WB Image

WB analysis of human epithelial carcinoma cells expressing FAK with (Lane 2-5) or without (Lane 1) 100 ng/mL Taxol for 18-24 hours in serum-free media using GTX24794 FAK (phospho Ser910) antibody. The data show that only the immunogen phosphopeptide blocks the signal, demonstrating the specificity of the antibody.



For full product information, images and publications, please visit our <u>website</u>.

Date 2025 / 12 / 27 Page 2 of 2