

À des fins de recherche uniquement

# Anticorps Monoclonal anti-XRCC5

Numéro de catalogue:**CL594-66546**



## Informations de base

Numéro de catalogue:	BC019027	Méthode de purification:
CL594-66546	Purification par protéine G	
Taille:	Identification du gène (NCBI):	CloneNo.:
100µl , Concentration: 908 µg/ml by Nanodrop;	7520	2G5E7
Hôte:	Nom complet:	Dilutions recommandées:
Mouse	X-ray repair complementing defective F 1:50-1:500	
Isotype:	repair in Chinese hamster cells 5 (double-strand-break rejoining)	Excitation/Emission maxima wavelengths:
IgG1	MW calculé	588 nm / 604 nm
Immunogen Catalog Number:	732 aa, 83 kDa	
AG9512	MW observés:	
	80-83 kDa	

## Applications

Applications testées:	Contrôles positifs:
FC (Intra), IF	IF : cellules HeLa,
Spécificité de l'espèce:	
Humain	

## Informations générales

There are at least two pathways for eukaryotes to repair DNA double-strand breaks: homologous recombination and nonhomologous end joining(NHEJ). The core NHEJ machinery includes XRCC4, DNA ligase IV and the DNA-dependent protein kinase complex, which consists of the DNA end-binding XRCC5/XRCC6 heterodimer and the catalytic subunit PRKDC. The heterodimer of XRCC5/XRCC6 enhanced the affinity of the catalytic subunit PRKDC to DNA by 100-fold. Once the XRCC5/6 dimer association with NAA15, it can bind to the osteocalcin promoter and activate osteocalcin expression. The XRCC5/6 dimer acts as a negative regulator of transcription when together with APEX1. Some published papers indicated that the MW of XRCC5 is 86kDa, while more papers suggested that XRCC5 is a 80kDa protein, as it was firstly introduced in publication. Thus, Ku80 and Ku86 are the same protein.

## Stockage

Stockage: Stocker à -20 °C. Éviter toute exposition à la lumière. Stable pendant un an après l'expédition.

Tampon de stockage:

PBS avec glycérol à 50 %, Proclin300 à 0,05 % et BSA à 0,5 %, pH 7,3.

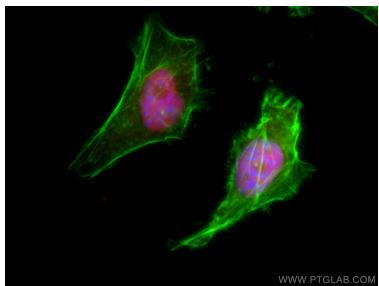
L'aliquotage n'est pas nécessaire pour le stockage à -20°C

\*\*\* Les 20ul contiennent 0,1% de BSA.

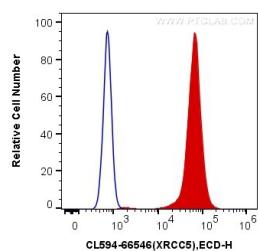
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## Données de validation sélectionnées



Immunofluorescent analysis of (4% PFA) fixed HeLa cells using Coralite®594-conjugated XRCC5 antibody (CL594-66546, Clone: 2G5E7) at dilution of 1:100.



1X10<sup>6</sup> HeLa cells were intracellularly stained with 0.2 ug Coralite®594 Anti-Human XRCC5 (CL594-66546, Clone:2G5E7) (red), or 0.2 ug Control Antibody. Cells were fixed and permeabilized with Transcription Factor Staining Buffer Kit (PF00011).