



## Aurora C (2 - 309)

Catalog Number (DU Number):  
**DU4009**

Accession:  
**ABO17332**

Expression  
**baculovirus**

Terminus and Tag:  
**N-Term 6His Uncleaved**

Purification Method:  
**Ni<sup>2+</sup>-NTA agarose**

Enzymatic Assay Format:  
**Standard filter binding assay**

Enzymatic Buffer:  
**50 mM Tris-HCl pH 7.5, 0.1 % 2-mercaptoethanol, 0.1 mM EGTA, 0.1 mM sodium vanadate, 10 mM magnesium acetate**

Enzymatic Substrate:  
**LRRLSLGLRRLSLGLRRLSLGL; Final concentration: 300  $\mu$ M**

Calculated Molecular Mass:  
**Mono-Isotopic Mass: 38,791.10**  
**Average Mass: 38,815.66**

Notes:  
**Following expression the culture is incubated with 50 nM okadaic acid for 1 hour prior to purification**

Purity:  
**>80 %**

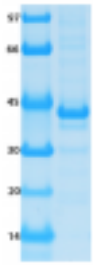
Storage Buffer:  
**50 mM Tris-HCl pH 8, 270 mM Sucrose, 150 mM NaCl, 0.1 mM EGTA, 0.1 % 2-mercaptoethanol, 0.2 mM PMSF, 1 mM Benzamidine.**

Storage Temperature:  
**-70 °C [Long term stability to be determined]**

Theoretical pI:  
**8.46**

Gel Information :

**Gel Image 1:**



Native Sequence:

**Amino acids S2 – S309 (end) of human Aurora C. Residue S29 of the fusion protein is equivalent to S2 of the native enzyme. The His(6) tag is located at residues 5 – 10. The following amino acid substitution is present: L – P, where L301 of the native enzyme is P328 of the fusion protein**

Protease Cleavage:

**rTEV (ENLYFQG) residues 18 - 24**

Cloning Sites:

**BamH1 and Sal1 sites of pFastBAC HTb**

Price per aliquot (100µg):

**£100.00**