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**Immobilized plasminogen domain**

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mAb		K4		K5-PD		BSA	
Competitor	Buffer	K4	Buffer	K5-PD	Buffer	K4	K5-PD
<b>Absorbance (OD<sub>405</sub>)</b>							
109	0.137±.003	0.137±.003	0.234±.012	0.227±.003	0.242±.01	0.127±.002	0.164±.003
51	<b>0.489 ±.019</b>	<b>0.309±.003</b>	0.293±.017	0.262±.002	0.241±.01	0.148±.001	0.214±.001
116	0.228±.032	0.182±.006	<b>0.908±.05</b>	<b>0.947±.03</b>	0.325±.02	0.180±.002	0.279±.01
Buffer	0.121±.004	0.122±.002	0.160±.009	0.146±.004	0.144±.004	0.113±.001	0.139±.002

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Binding of RIBS mAbs (290 nM) to immobilized isolated domains of Glu-plasminogen or to BSA, as a control, was determined in the presence of either buffer or soluble plasminogen domains (1 μM). Results are mean ± SEM; n= 3 for each experimental group. Bolded results indicate results that are significantly different (p<0.05) compared to corresponding controls with BSA immobilized onto the microtiter wells.