Table 7: Integrase

MAb ID	HXB2 Location	<b>Author's Location</b>	Sequence	Neutralizing	Immunogen	Species(Isotype)		
166 1C4	Integrase(1–16)	Integrase(1–16 HXB2)	FLDGIDKAQDEHEKYH	no	bacterial expressed integrase	murine( $IgG_1\kappa$ )		
	<b>References:</b> [Haugan (1995), Nilsen (1996)]							
		feres with integrase binding to	<u> </u>					
		•	vith the N-terminal part of inte	-				
	6G5, 7B6, 7C6 - -Nilsen96	- these MAbs inhibit end prod	cessing and DNA joining, but	had little effect	on integration activities	3		
167 2C11	Integrase(1–16)	Integrase(1_16 HXR2)	FLDGIDKAQDEHEKYH	no	bacterial expressed	murine( $\operatorname{IgG}_1 \kappa$ )		
107 2011	integrase(1 10)	micgrase(1 10 11/102)	TEDOIDKAQDEHEKTH	110	integrase	marme(1gO <sub>1</sub> n)		
	References: [Nilsen	(1996)]			•			
	• 2C11: One of a la	arge set of MAbs that interact	with the N-terminal part of inte	egrase: 1C4, 2C1	1, 2E3, 3E11, 3F9, 5F8	,		
	6G5, 7B6, 7C6 -	these MAbs inhibit end prod	cessing and DNA joining, but	had little effect	on integration activities	3		
	–Nilsen96							
168 2E3	Integrase(1–16)	Integrase(1–16 HXB2)	FLDGIDKAQDEHEKYH	no	bacterial expressed integrase	$murine(IgG_1\kappa)$		
	<b>References:</b> [Nilsen (1996), Ovod (1992)]							
	• 2E3: There are two MAbs called 2E3 – the other one binds to Nef –Ovod92							
	• 2E3: One of a large set of MAbs that interact with the N-terminal part of integrase: 1C4, 2C11, 2E3, 3E11, 3F9, 5F8,							
	6G5, 7B6, 7C6 - -Nilsen96	- these MAbs inhibit end prod	cessing and DNA joining, but	had little effect	on integration activities	S		
169 3E11	Integrase(1–16)	Integrase(1–16 HXB2)	FLDGIDKAQDEHEKYH	no	bacterial expressed integrase	$murine(IgG_1\kappa)$		
	References: [Otteken (1992), Nilsen (1996)]							
	• 3E11: There is another MAb with this ID that recognizes p17 –Otteken92							
	• 3E11: Recognized an epitope present on HIV-2/SIVmac, SIVagm, HIV-1, and SIVmnd –Otteken92							
	• 3E11: One of a large set of MAbs that interact with the N-terminal part of integrase: 1C4, 2C11, 2E3, 3E11, 3F9, 5F8,							
	6G5, 7B6, 7C6 – these MAbs inhibit end processing and DNA joining, but had little effect on integration activities							
	6G5, 7B6, 7C6 -	<ul> <li>these MAbs inhibit end prod</li> </ul>	cessing and DNA joining, but	had little effect	on integration activities	3		

MAb ID	<b>HXB2</b> Location	Author's Location	Sequence	Neutralizing	Immunogen	Species(Isotype)			
170 3F9	Integrase(1–16)	Integrase(1–16 HXB2)	FLDGIDKAQDEHEKYH	no	bacterial expressed integrase	murine( $\operatorname{IgG}_1 \kappa$ )			
	References: [Nilsen	· -	with the N terminal part of inte	grass, 1C4, 2C1	1 202 2011 200 500				
		• 3F9: One of a large set of MAbs that interact with the N-terminal part of integrase: 1C4, 2C11, 2E3, 3E11, 3F9, 5F8, 6G5, 7B6, 7C6 – these MAbs inhibit end processing and DNA joining, but had little effect on integration activities –Nilsen96							
171 5F8	Integrase(1–16)	Integrase(1–16 HXB2)	FLDGIDKAQDEHEKYH	no	bacterial expressed integrase	$murine(IgG_1\kappa)$			
	<ul> <li>References: [Haugan (1995), Nilsen (1996)]</li> <li>5F8: There is another MAb with this ID that recognizes and unknown protein in HIV –Pinter95b</li> <li>5F8: MAb interferes with integrase binding to DNA –Haugan95</li> <li>5F8: One of a large set of MAbs that interact with the N-terminal part of integrase: 1C4, 2C11, 2E3, 3E11, 3F9, 5F8, 6G5, 7B6, 7C6 – these MAbs inhibit end processing and DNA joining, but had little effect on integration activities –Nilsen96</li> </ul>								
172 6G5	Integrase(1–16)	Integrase(1–16 HXB2)	FLDGIDKAQDEHEKYH	no	bacterial expressed integrase	$murine(IgG_1\kappa)$			
	References: [Nilsen (1996)]  • 6G5: One of a large set of MAbs that interact with the N-terminal part of integrase: 1C4, 2C11, 2E3, 3E11, 3F9, 5F8, 6G5, 7B6, 7C6 – these MAbs inhibit end processing and DNA joining, but had little effect on integration activities –Nilsen96								
173 7B6	Integrase(1–16)	Integrase(1–16 HXB2)	FLDGIDKAQDEHEKYH	no	bacterial expressed integrase	$murine(IgG_1\kappa)$			
	References: [Nilsen (1996)]  • 7B6: One of a large set of MAbs that interact with the N-terminal part of integrase: 1C4, 2C11, 2E3, 3E11, 3F9, 5F8, 6G5, 7B6, 7C6 – these MAbs inhibit end processing and DNA joining, but had little effect on integration activities –Nilsen96								
174 7C6	Integrase(1–16)	Integrase(1–16 HXB2)	FLDGIDKAQDEHEKYH	no	bacterial expressed integrase	$murine(IgG_1\kappa)$			
	References: [Nilsen (1996)]  • 7C6: One of a large set of MAbs that interact with the N-terminal part of integrase: 1C4, 2C11, 2E3, 3E11, 3F9, 5F8, 6G5, 7B6, 7C6 – these MAbs inhibit end processing and DNA joining, but had little effect on integration activities –Nilsen96								

## **HIV Monoclonal Antibodies**

MAb ID	<b>HXB2</b> Location	Author's Location	Sequence	Neutralizing	Immunogen	Species(Isotype)		
175 6C5	Integrase(17–38)	Integrase(17–38 HXB2)	SNWRAMASDFNLPPVVA KEIVA	A- no	bacterial expressed integrase	murine( $\operatorname{IgG}_1 \kappa$ )		
	• 6C5: MAb interfe	References: [Haugan (1995), Nilsen (1996)]  • 6C5: MAb interferes with integrase binding to DNA –Haugan95  • 6C5: This MAb inhibits end processing and DNA joining, but had little effect on integration activities –Nilsen96						
176 4D6	Integrase(42–55)	Integrase(42–55 HXB2)	KCQLKGEAMHGQVD	no	bacterial expressed integrase	murine( $\operatorname{IgG}_1\kappa$ )		
	• 4D6: MAb interfe	(1995), Nilsen (1996)] eres with integrase binding to nhibits end processing and Di	DNA –Haugan95 NA joining, and reduces reinteg	gration activity -	-Nilsen96			
77 anti-K159	Integrase(151–163)	Integrase(163–175)	VESMNKELKKIIG		peptide K159 of Integrase	rabbit(IgG)		
	<ul> <li>References: [Maroun (1999)]</li> <li>anti-K159: Both the peptide K159, SQGVVESMNKELKKIIGQVRDQAEHLKTA, and the Abs raised against this peptide inhibit Integrase activity – K159 was found to fulfill condition of minimal number of helical heptads to achieve the formation of a stable coiled-coil structure – Integrase is proposed to function as a dimer interacting in this region –Maroun99</li> </ul>							
178 8-6	Integrase(211–227)	Integrase(211–227 HXB2	) KELQKQITKIQNFRVYY	no	Integrase linked to maltose binding protein (MBP)	murine(IgG <sub>1</sub> )		
	<ul> <li>Donor: Yoshihiro Kitamura, Div of Mol Genetics, Nat Inst of Infectious Diseases, Musashimurayama, Japan</li> <li>References: [Ishikawa (1999)]</li> <li>8-6: Antibody binds proximal to the DNA binding region –Ishikawa99</li> </ul>							
179 2-19	Integrase(228–236)	Integrase(228–236 HXB2	) RDSRNPLWK	no	Integrase linked to maltose binding protein (MBP)	murine(IgG <sub>2b</sub> )		
	<ul> <li>Donor: Yoshihiro Kitamura, Div of Mol Genetics, Nat Inst of Infectious Diseases, Musashimurayama, Japan</li> <li>References: [Ishikawa (1999)]</li> <li>2-19: MAb inhibits RT-Integrase interaction, and the terminal cleavage and strand transfer functions of Integrase, but not the disintegration activity –Ishikawa99</li> </ul>							

MAb ID	<b>HXB2</b> Location	Author's Location	Sequence	Neutralizing	Immunogen	Species(Isotype)		
180 19	<ul> <li>19: BALB/c mice and the antibodies</li> <li>19: scAb2-19: A binding to integra</li> </ul>	Integrase(228–236 LAI) Bender (1994), Levy-Mintz (19 e were immunized with rec in s characterized – 19 has a low a single-chain Ab made from se – scAb interfered with the fed cells, but the virus produc Kitamura99	ntegrase, hybridomas express binding affinity –Bizub-Ben MAb 2–19 –acts intra-cellu folding of Gag-Pol polyprote	der94 larly to block inf in, the Ab did not	ection at low MOI by affect viral production	$murine(IgG_1)$		
181 8-22	References: [Ishikaw	maltose binding protein (MBP)  Donor: Yoshihiro Kitamura, Div of Mol Genetics, Nat Inst of Infectious Diseases, Musashimurayama, Japan  References: [Ishikawa (1999)]  • 8-22: MAb inhibits the terminal cleavage and strand transfer functions of Integrase, but not the disintegration activity						
182 4-20	Integrase(253–261)	Integrase(253–261 HXB2	) DNSDIKVVP	no	Integrase linked to maltose binding protein (MBP)	murine(IgG <sub>1</sub> )		
	<ul> <li>Donor: Yoshihiro Kitamura, Div of Mol Genetics, Nat Inst of Infectious Diseases, Musashimurayama, Japan</li> <li>References: [Ishikawa (1999)]</li> <li>4-20: Inhibits the terminal cleavage and strand transfer functions of Integrase, but not the disintegration activity         <ul> <li>Ishikawa99</li> </ul> </li> </ul>							
183 6-19	Integrase(261–270)	Integrase(261–270 HXB2)	) RRKAKIIRD	no	Integrase linked to maltose binding protein (MBP)	$murine(IgG_{2b})$		
	<ul> <li>Donor: Yoshihiro Kitamura, Div of Mol Genetics, Nat Inst of Infectious Diseases, Musashimurayama, Japan</li> <li>References: [Ishikawa (1999)]</li> <li>6-19: Inhibits the terminal cleavage and strand transfer functions of Integrase, but not the disintegration activity         <ul> <li>Ishikawa99</li> </ul> </li> </ul>							

## **HIV Monoclonal Antibodies**

MAb II	HXB2 Location	<b>Author's Location</b>	Sequence	Neutralizing	Immunogen	Species(Isotype)			
184 8E5	Integrase(262–271)	Integrase(262–271 HXB2)	RRKAKIIRDY	no	bacterial expressed integrase	$murine(IgG_1\kappa)$			
	References: [Haugan	(1995), Nilsen (1996)]			C				
		• 8E5: MAb interferes with integrase binding to DNA –Haugan95							
		ree MAbs recognize an epitope IN – these MAbs inhibit end- lilsen96							
185 7C3	Integrase(262–271)	Integrase(262–271 HXB2)	RRKAKIIRDY	no	bacterial expressed integrase	murine( $\operatorname{IgG}_1 \kappa$ )			
		n (1995), Nilsen (1996)]							
		Feres with integrase binding to $\Gamma$							
	react with HIV-2	<ul> <li>7C3: A set of three MAbs recognize an epitope in this region, 7C3, 7F11, and 8E5 – all three HIV-1 MAbs cross-react with HIV-2 IN – these MAbs inhibit end-processing, DNA joining and reintegration, and had little effect on disintegration –Nilsen96</li> </ul>							
186 7F11	Integrase(262–271)	Integrase(262–271 HXB2)	RRKAKIIRDY	no	bacterial expressed integrase	$murine(IgG_1\kappa)$			
	References: [Nilsen (	<b>References:</b> [Nilsen (1996), Lasky (1987)]							
	<ul> <li>7F11: A set of th react with HIV-2 disintegration –N</li> </ul>	<ul> <li>7F11: A set of three MAbs recognize an epitope in this region, 7C3, 7F11, and 8E5 – all three HIV-1 MAbs cross-react with HIV-2 IN – these MAbs inhibit end-processing, DNA joining and reintegration, and had little effect on disintegration –Nilsen96</li> <li>7F11: There is another MAb with this name that binds to gp120 –Lasky87</li> </ul>							
187 MAb 35	Integrase(264–273) References: [Barsov	Integrase() (1996), Acel (1998)]	KAKIIRDYGK	no	rec IN	murine( $\operatorname{Ig}G\kappa$ )			
	<ul> <li>MAb 35: There a</li> <li>MAb 35: Althoundisintegration –B</li> <li>MAb 35: Integra</li> </ul>	<ul> <li>MAb 35: There appears to be two different IN Abs with similar names: MAb 35 and 35 –Barsov96,Bizub-Bender94</li> <li>MAb 35: Although MAb 35 does not inhibit HIV-1 IN, Fab 35 inhibits 3'-end processing, strand transfer and disintegration –Barsov96</li> <li>MAb 35: Integrase was shown to have intrinsic DNA polymerase activity that can catalyze gap repair – MAb 35 inhibits this activity –Acel98</li> </ul>							