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#### (54) MONOCLONAL ANTIBODIES AGAINST TISSUE FACTOR PATHWAY INHIBITOR

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#### (57) ABSTRACT

Isolated monoclonal antibodies that bind human tissue factor pathway inhibitor (TFPI) and the isolated nucleic acid molecules encoding them are provided. Pharmaceutical compositions comprising the anti-TFPI monoclonal antibodies and methods of treating deficiencies or defects in coagulation by administration of the antibodies are also provided. Methods of producing the antibodies are also provided.

Figure 1

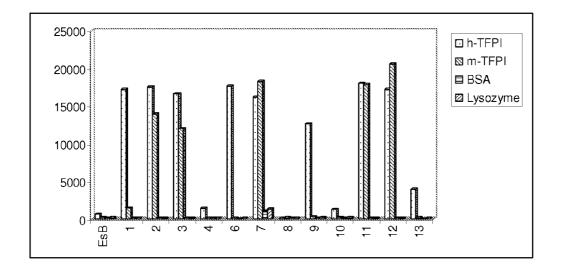


Figure 2

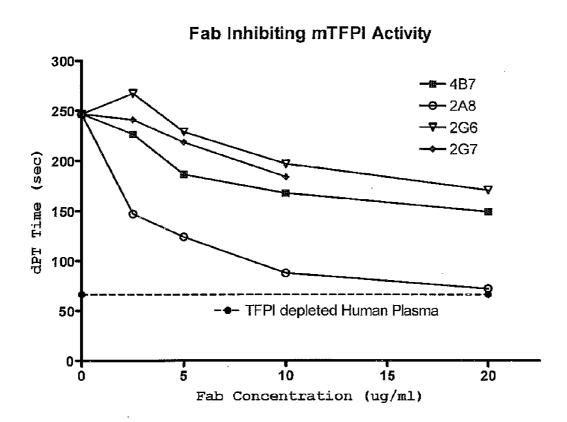


Figure 3

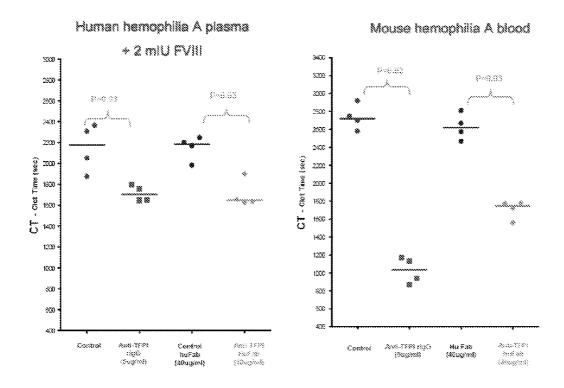


Figure 4

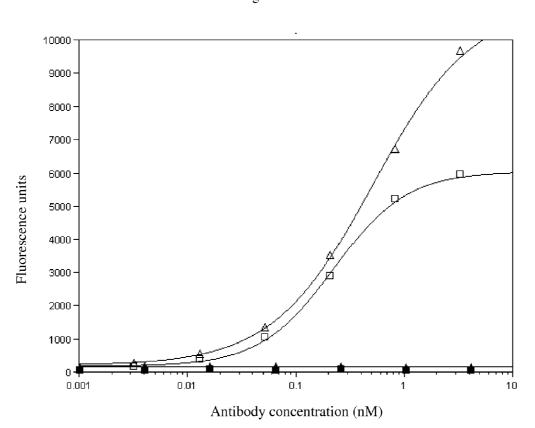
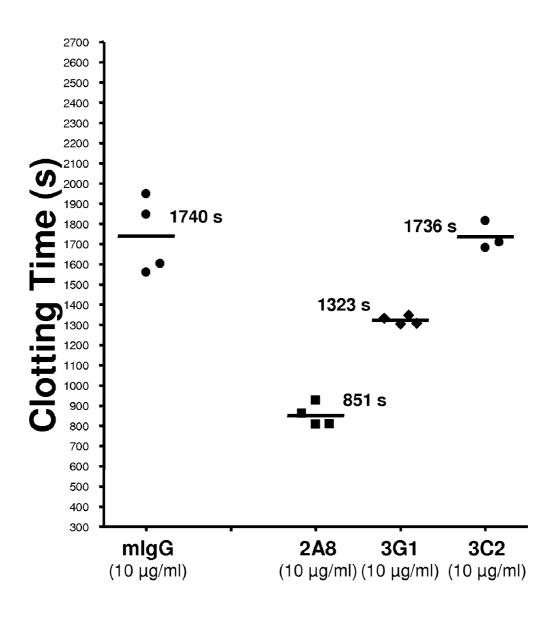


Figure 5



Figure

TP-2A10 DIELIQ-PPSVSVAPGQIARISCSGDKLGK TP-2B1 DIELIQ-PPSVSVAPGQTARISCSGDNLGN	KYVHWYQQKPGQAPVLVIYGDDKRPSGIPERFSGSNSGNIATLTISSTQABDEADYYCQAWGSISRFVFGGGTK-TVLGQ KVAHWYQQKPQQAPVIVIYYDNKRPSGIPERFSGSNSGNIATLTISSTQAEDRADYYCQSWTPGS-NTPVFGGGTR'TVIGQ
	EGYVFGGGTK
TE-ZAS LIELIÇ-FESVSVAPGÇIAKISCSGDNERN	VFVFGGGTK-
II ZBS DIELIQ IISVSVANGQIAKISCSGDNEKG IP-2G7 DIELIG-PPSVSVAPGOIAKISCSGDNLGI	IIASWIQQALYOJAIYIIEENNALSEIFEKK SESNSENIAILIISSIQAEDEADIISEGONDSIYI YHVEGGEIKLIYLGO YYYHWYOOKPCOAPVIVIYEENNKESEIPEKFSESNSENIAILIISSIOAEDEADYYCOIYDSNN-ESIVFECIKLIYLGO
TP-2G4 DIELIQ-FPSVSVAPGQTARISCSGDALRK	
	NA-VFGGGTK
	YYAHWYQQKPGQAPVLVIYYBSKRFSGIPERFSGSXSGNIYILTISGTQAEDEADYYGQS-ISKVFGGGTK-TVLGQ
TP-3EL DIEDIQ-FESVSVAPGQTAKISCSGDNIGS TP-3El DIEDIG-PESVSVAPGGTARISCSGDNIGS	YYANWYQQKEGQAFVLVIYDDSNKESGIFEKESGSNSGNIAILIISGIQAEDEADYYCGSYD-SI-GLLVEGGGIK-IVLGQ YPASWYOOKPGQAPVIVIYDDSNEPSGIPERFSGSNSGNIAILIISGIOAEDRADYYCEGSNVFGGCIK'IVIGO
DIELTO-PPSVSVAPGOTA	TYTSRS-HSYVFGGGTK7
II 4A7 DIELIQ PPSVAPGQTARISCSGDALGS	
TP-4G8 DIELTQ-FPSVSVAPGQIAZISCSGDKLGS	KSVHWYQQKPGQAPVLVIYRDIDRFSGIPERFSGSNSGNIAILISGTQAEDEADYYCQIYDYILN-VFGGGTK_IVLGQ
TP-2B3 DIVLIQSPATISLSPGERATLSCRASQNIGSN-	YLAWYQQKPGQAPRLLIYGASTRATGVPARFNGSGSGTDFTLTISSLEPEDFAVYYQQQLNSIPVTFGQGTKVFIRRT
TP-2F9 DIVLIQSPATISLSPGERATLSCRASQSVSSQ-	YLAWYQQKPGQAPRLLIYAASSRATGVPARFSGSGSGTDFTLTISSLEPEDFAVYYQQQSNLPATFGQGTKVFIKRT
TP-2G5 DIVLIQSPATISLSPGBRATLSCRASQNVSSN-	YLAWYQQKPGQAPRLLIYDASNRATGVPARFSGSGSGTDFTLTISSLEPEDFAVYYCQQFYDSPÇTFGÇGTKVEIKRT
DIVLIQSPATISLSPGER	-
TP-2B9 DIÇMIQSFSSJSASVGDRVIITCRASQSIRS	YLAWYQQKPGKAPKLLIYKASNIQSGVPSRFSGSGSGTDFTLTISSLQPEDFAVYYCHQYSDSPVTFGQGTKVEIKRT
DIÇMTQSFSSLSASVGDRV	-
DIÇMTQSFSS_SASVGJRV	
DIÇMTQSPSSLSASVGDRV	
DIOMTOSESSISASVGDRA	-
DIÇMTÇSPSSLSASVGDRV	-
DIÇMTQSPSSLSASVGDRV	
DIÇMTQSFSSISASVGDRV	-
TP-2D7 DIVMIQSP S'PVTPGRPASISCRSSQSLLHSN	STSCRSSQSLIHSNGYTVISWYIQKPGQSPQILITYIGSNRAGGVPDRFSGSGSGTDFTIKTSRVRARDVGVVVCQQYDNAPTTFGGGTKVFTKRT
DIVETOSP-S-PVTPG3-29	KSISCRSSQSLVESDGNITLINWTLQKPGQSPQLLIYKGSNRASGVPDRESGSGSTDFILKISRVEAEDVGVYYQQYDSYP LIFGGCTKVEIKRT
DIALTQ-PASVSGSPGQSITISCTGTSSDIGGY	NYVSWYQQHPGKAPKLMIYGVNYFPSCVSNFFSGSKSGNIASLTISGLQAEDEADYYGSSADKFT-MSIVFGGCIK-IVLGQ
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DIALIQ-PASVSGSPGQSI	NTVSWYQQHPGKAPKLMIYSVSSRPSGVSNRFSGSKSGNIASLTISGLQAEDEADYYCQSYDLNNLVFGGGTK-TVLGQ
IP-3A4 DIELIQ-PESVSVAPGQIARISCSGDNLRD	KYASWYQQKPGQAPVLVIYSKSEKPSGIPERFSGSNSGNIATLTISGTQAEDEADYYCSSYTLNPNLNYVFGGGTKITVLGQ
	-VKFAVFGGGTK_
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	TRSLVFGGGTKI
	KYVSWYQQKPGQAPVLVIYGDNKRFSGIPERFSGSNSGNIAILISSTQAEDEADYYCQSYTYSLNÇVFGGGTK_TVLGQ
TP-3B3 DIELIQ-PASVSVAPGQIARISCSGDNLRS	KYAHWYQQKPGQAPVLVIYGDNNKFSGIPERFSGSNSGNIATLTISGTQAEDEADYYCSAYAMGSSPVFGGGTKITVLGQ

	SOGTLVTVSS	SQUILVIVSS	-YYYGFDYWGQGTLVTVSS	THFDVWGQGTLVTVSS	SYLGYFDVWGQGTLVTVSS	XYRYWEDYWGQGTLVTVSS	-G-PGMDVWGQGTLVTVSS	-HMDYWGQGTLVTVSS	MDSWGQGTLVTVSS	-GDIYFDYWGQGTLVTVSS	-GYFDYWGQGTLVTVSS	-SSDYWGQGTI,VTVSS	GGFDYWGQGTLVTVSS	TCFDYWCQCTLVTVSS	-WGFDYWGQGTLVTVSS	GFDVWGQGTLVTVSS	S-GVMDVWGQGTT,VTVSS	NGTDVWGQGTLVTVSS	LYVYSDVWGQGTLVTVSS	YYAYVDIWGQGTLVTVSS	YFDYWGQGTLVTVSS	SQCTL,VTVSS	SOGILVIVSS	SQUILVIVSS	SQUILVIVSS	SQUILVIVES	SQCTT,VTVSS	SQUILLVIVSS	MHYKGMDIWGQGTLVTVSS	GLIDVWGQGTLVTVSS	SOGILVIVES	SQCTLATVSS	SQUILVIVSS	SOCILVIVSS	SOGILVIVSS	FYFDVWGQGTLVTVSS	NDSCWFDVWCQCTLVTVSS	SQUILVIVSS	RYSLGADSWGQGTLVTVSS	YYCKCVDLWGQGTLVTVSS	YGGMDYWGQGTLVTVSS	GGGVDYWGQGTLVTVSS	-GGGVDYMGQGTLVTVSS	ココネエネコエラブに
33	-YKESYFDIWGQGTLVTVSS	-LFPGYFDYWGQGTLVTVSS	YYYYGEDYW	THEDVW	-SYLGYFDVW	-XYRYWEDYW	G-PGMDVW	HMDYW	MDSW	GDIYF'DYWC	GYFDYW	SSDYW	GGF'DYWC	ICEDYW	MGF.D.X.MC	GEDVWC	S-GVMDVW	NGTDVWC	LYVYSDVW	YYAYVDIW	YFDYW	-XYDSGFDVWGQGTLVTVSS	SGFTFNNNAISWVRQA-GKGLEWVSAINS-SSSSTSYADSVKGKFTISKINSKNTLYIQMNSLRAEDTAVYYCARGHRRGHSWASFIDYWGQGTLVTVSS	TIGVLWDDVWGQGTLVTVSS	-LPYMVFDYWGQGTLVTVSS	-YPYLVFAIWGQGTLVTVSS	GYCHYYPFDYWGQGTT,VTVSS	-GDSYMYDVWGQGTLVTVSS	MHYKGMDIW	GLIDVW	-GIIGFFFSDIWGQGTLVTVSS	-VYYGFDFWGQGTT,VTVSS	-GWIYSYIDVWGQGILLVIVSS	==SZZNSKNTLYZQMNSLRAEDTAVYYCARPIKAGZIWWWGPYMDVWGQGTLVTVSS	YPOWGWYTDVWGQGTLVTVSS	FYFDVW	-NDSCWEDVWC	-YMNLLAG-WGQGTLVTVSS	RYSLGADSWO	-YYCKCVDLW	YGGMDYW	GGGVDYW	GGGVDYW	יייייטעוניוקט ז א־
CUKS	H	İ	ARVG	ARVNIS	AR	AR	ARIMSKYG-	ARYSSIG	ARQDGYGG	ARWEG_AD-	ARSNWS	ARVNQV"	AKAKAKAS	ARHNPD	AKWHSUKA	ARIISMV	ARGG-	AREGULNA-	ARYLGSN	ARVNANG	ARGVHS	1	ARCHHRG	AR	ARAD	ARGDY	ł		AR	ARNG	İ	-	AKQAG(	ARPIKAGRI		AKYDN	AR	AK12	ARYW	AR	AK	ARAL	ARAI	AKINGK
	RAEDTAVYYC	RAEDTAVYYC	RAEDTAVYYC	RAEDTAVYYC	RAEDTAVYYC	RAEDTAVYYC	TPEDIAVYYC	TPEDIAVYYC	TPEDTAVYYC	TPEDTAVYYC	TPEDIAVYY	TPEDTAVYYC	TPEDTAVYYC	TPEDTAVYY	TPEDTAVYYC	TPEDIAVYY	TPEDTAVYYC	TPEDTAVYYC	TPEDTAVYY	TPEDTAVYYC	RSEDTAVYYC	KASDTAMYYC	RAEDTAVYYC	RAEDTAVYY	RAEDTAVYYC	RAEDTAVYYC	RAEDTAVYYC	RAEDTAVYYC	RAEDTAVYYC	RAEDTAVYYC	RAEDTAVYY	RAEDTAVYYC	RAEDTAVYYC	RAEDTAVYY	RAEDTAVYYC	RAEDTAVYY	RAEDTAVYY	RAEDTAVYYC	RAEDTAVYYC	RAEDTAVYYC	RAEDTAVYY	RAEDTAVYY	RAEDTAVYYC	KALDIRVII
	NTLYZQMNSL	==SSCNSKNTLY_QMNSLRAEDTAVYYCARG	NILYZOMNSL	<pre>====================================</pre>	SRONSKNTLYLQMNSLRAEDTAVYYCAR	NILYLOMNSL	NOFSLOLNSV	NOFSLOLNSV	NQFSLQLNSV	NQF'S_Q_NSV	NQFSZQZNSV	""SKNQFS",Q",NSVTPRDAVYYCARVNÇ	NFJTSKNQFSLQLNSVTPEDTAVYYCARARAS	NQFSLQLNSV	NFJTSKNQFS_Q_NSVTPEDTAVYYCARWHSDKA	NOFSZQZNSV	ASKNOFSTOTASVTPFIDTAVYYCAR	NQF'S_Q_NSV	NOFSLOLNSV	NOFSLOLNSV	STAYMELSSL	STAY QWSST	NTLYLOMNSL	NTLYLOMNSL	NTLYLOMNSL	TISRONSKNILYLQMNSLRAEDIAVYYCARGDY	-P TASKNITI-YOMNST.RAEDTAVYYCART M-	SRUNSKNTLY QMNSLRAEDTAVYYCAR	TISRINSKNILYIQMNSLRAEDTAVYYCAR	NILYZOMNSL	<b>==SRCNSKNTLYLQMNSLRAEDTAVYYCARKGG</b>	SRNTTYTOMNSTRAEDTAVYYCAR-	SACNSKNTLYLQMNSLRAEDTAVYYCARQAG-	NILYLOMNSL	SRONSKNTLYOMNSLRAEDTAVYYCAR-	NILYLOMNSL	NILYZOMNSL	SKUNSKNTLY JOMNSTRAEDTAVYYCAKLY	NILYZOMNSL	NILYZOMNSL	NILYLOMNSL	NITYZOMNSI	SACNSKNTLYLOMNSLRAEDTAVYYCARAL 	NILIIV
	WELESRONSKI	WEISSONSKI	WIISKINSK	WIISKONSK	RIISKINSKI	WIISRONSKI	NSICENIES	METCENTER	MSICONIES	ASTC-N	MSICENIER	RTTCENS	ASTUS NOTES IN	STINDIEN SKI	ASTUSKI	MSTCENTER	1	ASTUS NO SERVICE	NSTCCNIISK	MSICENIER	VISICEIXIA	V-SACKSTV	WIISKINSKI	WIISKINSK	WEISSCNSKI	WIISKINSK	i.	ASKLASAL BY	TISKINSKI	WIISKINSKI	WIISKONSKI	RTTSRVCK	A SACNSKI	WIISKONSKI	TISKINSKI	WIISKINSK	WIISKONSKI	RSKUSK	TISKINSKI	TISKINSKI	WIISKONSK	TISKINSKI	TISSONSK	Man-y-a II
	TILYADSVKGI	TNYADSVKGE	ITYYADSVKCE	TYYADSVKCE	TYYADSVKCE	TYYADSVKCI	TNSYAVSVKSE	TNDYAVSVKSE	TNSYAVSVKSE	INSYAVSVKSE	TNITYAVSVKSI	NDYAVSVKSE	NUYAVSVKSE	NDYAVSVKSE	INDYAVSVKS	TNDYAVSVKSI	TNEYAVSVKSE	INDYAVSVKSE	TNAYAVSVKSE	INDYAVSVKSE	TKYAQKFQGE	TNYSPSFQC(	TSYADSVKCE	TYYADSVKCE	TYYADSVKCE	THYADSVKG	TYYADSVKCF	TYYADSVKCE	SSSNTYYGDSVKGRF	TYYADSVKGE	ITYYADSVKGI	ITNYADSVKCE	TYYADSVKG	ITYYADSVKGI	TSYADSVKG	TEYADSVKG	ITYYADSVKCE	TYYADSVKG	TNYADSVKG	TYYADSVKG	TYYADSVKCE	TYYADSVKG	TYYADSVKGE	1 I TRUDVING
CDR2	'SAISY-TGSI	'SNISY-MGSI	'SNISY-SGSI	SVISG SGS	'S_ISG-VSS	SSIRG-SSS	GMIYYRSKW	GMIYHRSKW	GFIYRRSKW	GHIYYRSKW	GIIYYRSKW	GMTYYRSKW	GVIYYRSKW	CFIYYRSKW	GLIYKRSKW	GIIYYRSKW	GMTYYRSKW	GLIYYRSKW	GLIYYRSKW	GMIFYRSKW	GTIFF-YDG1	ICTIQP-SDSI	'SAINS-SSS	SIES VSS	'SVISG-SSS	'SVIEY-SGSF	7SSTTS-SSF	SSTSS-SSTSS,	SSISS SSSI	'SSIKG-SGS	SNISS-NSSI	'SNTSS-SGSI	STISY-DGSE	SVISS-VGSP	'SAILS-DGS	'SGISG-DGSL	'SAIKS-DCSN	SGISY-NGS	SGISY NGS	'SGIQY-DGS)	SSISY-DSSN	SGISY SSS	'SGISY-SSSE	מדדמפוופפד
	CAPGKGLEWN	CAPGKGLEWV	ÇA∋GKGLEWV	QAPGKGLEWV	CAPGKGLEWV	CAPGKGLEWN	CSPGRGLEWI	CSFGRGLEWI	ÇSPGRELEWI	TMHTDYB/SÖ	QSPGRGLEWI	CSPGRGI,FWI	TMHTDYDESÖ	ÇSPGRGLEWI	TMHTDY:DESÖ	QSPGRGLEWI	QSPGRGT.RWI	TMHTDYB/SÖ	QSPGRGLEWI	CSPGRGLEWI	CAPGOGLEWN	QMPGKGT,FWW	CAPGKGLEWV	CAPGKGLEW	CAPGAGLEW	CAFGKGLEWN	QA⊃GKGT.FWV	CAPGAGLEWV	QAPGKGLEWV	CASGKGLEW	CAPGKGLEW	QA>GKGT.FWV	ÇAZGAGLEWV	CAPGKGLEWN	<u>ÇAFGKGLEW</u>	CAPGAGLEWN	QAPCKCLEWV	ÇAZGAGLEWV	ÇAZGKGLEWN	CAPGAGLEW	ÇAFGKGLEWV	ÇAZGKGLEWN	CAPGKGLEWN	LASTGAGLEWY
CURI	SGFTFTSYSMYWVRÇA=GXGLEWVSAISY-TGSNTLYADSVKGRFTISCNSKNTLYIQMNSLRAEDTAVYYCARAFLG	SGFTFSSYCMHWVRQAPGKGLEWVSNISY-MGSNTNYADSVKGRF	FTFSSYAMIWVRÇAFGKGLEWVSNISY-SGSNTYYADSVKGKFTTSKUNSKNTLYLQMNSLRAEDTAVYYCARVG	FIFSSYGS WVRQAPGKGLEWVSVISG SGSSTYYADSVKGRF	SGFTFSSYGMSWVRQAPGKGLEWVSLISG-VSSSTYYADSVKGRE	SGFTFRSYSWSWVRQA-GKGLEWVSSIRG-SSSTYYADSVKCRFTTSRNSKNTLYLQMNSLRAEDTAVYYCAR	SGDSVSSNSAAWGWIRQSFGRGLEWLGMIYYRSKWYNSYAVSVKSRIIINFITSKNQFSLQINSVIPEDIAVYYCARIYSKYG	SGDSVSSNSAAWGW_RÇSFGRGLEWLGM1YHRSKWYNDYAVSVKSRNFJTSRNQFSLQ_NSVTFEDTAVYYCARYSSTG	SGDSVSSNGAAWGWIRQSJGRGLEWLGFIYRRSKWYNSYAVSVKSRIIINJISKNQFSLQINSVITBDIAVYYCARQDGXGG	SGDSVSSNGAAWGW_RQS-GGGLEWLGHLYYRSKWYNSYAVSVKSRN-JTSKNQFS_Q_NSVTFEDTAVYYCARWGG_AD	SGDSVSSNSAAWGWIRQSFGRGLEWLGIIYYRSKWYNIYAVSVKSRIIINFFITSNFFKNQFSLQINSVTPEDTAVYYCARSNWS	SGDSVSSNSAAMGW_RQSPGRGI,FWI,GMTYYRSKWYNDYAVSVKSR_	SGDSVSSNSAAWEW_RÇSFGRGLEWLGV1YYRSKWYNDYAVSVKSK_	SCDSVSSNSAAMSWIRÇSPGRGLEWLGFIYYRSKWYNDYAVSVKSRIIINPITSNRNQFSLQINSVTPEDTAVYYCARHNPD-	SGDSVSSNSAAWSW_RQS=GRGLEWLG11YKRSKWYNDYAVSVKSR_	SGDSVSSSSAAMSWIRÇSFG7GLEWLG11YYRSKWYNDYAVSVKSRIIINFOTSKNQFSLQLNSVTPEDTAVYYCARHSYV-	SCDSVSSSSAAWSW_RQS>GRGLFWLGMTYYRSKWYNFYAVSVKSR_	SGDSVSSNSAAWGW_RQS=GAGLEWLG_LYYRSKWYNDYAVSVKSRN=JW=JWSKNQFS_Q_NSVTPEDWAVYYCARE GD_NA-	SGDSVSSNSGEWGWIRQSJGGGLEWLGITYYRSKWYNAYAVSVKSRITTNJJTSKNQFSLQINSVTPEDTAVYYCARYIGSN	SGDSVSSNSAAWSW_RQSFGRGLEWLGMIFYRSKWNNDYAVSVKSRNPTTSKNQFS_Q_NSVTPEDTAVYYCARVNANG	SGYTFTGNSMHWVRÇAFGQGLEWMGTIFP-YDGTTKYAQKFQGRVIXIRJTSISTAYMELSSLRSEDTAVYYCARGVHS-	SGYSPTDYWTCWVR@V>GKGLRWWGTTQP-SDSDTNYSPSFQCQV <sup></sup> SAJKSTSTAYJQWSS1.KASDTAMYYCARFYWWG-	FNNNAISWVF	FTFNDYAMSWVRQA-GKGLEWVSLIBS VSSSTYYADSVKGRFTISRNSKNTLYLQMNSLRAEDTAVYYCAR	FTFRNYAMNWVRQAEGKGLEWVSVISG-SSSYTYYADSVKGRFT.ISRNJAMNUNTYLQMNSLRAEDTAVYYCARAD	FTFSDEAMHWVRQAFGKGLEWVSVIEY-SGSKTNYADSVKGRF	PTFSRYAMSWVRQAPGKGLEWVSSTIS-SSSETYYADSVKCRF	FTFSPYVMSWVRQA-GKGLEWVSSLSS-SSSNTYYADSVKGRE	FIFSSYSMVRQAPGKGLEWVSSISS	FTFSSYSMSWVRQASGKGLEWVSSIKG-SGSNTYYADSVKCRF   ISRINSKNTLYIQMNSLRAEDTAVYYCARNC	TTINSYAMSWVRÇAFGKGLEWVSNISS-NSSNTYYADSVKGRF	FTFNSYYMSWVRQAPGKGLFWVSNTSS-SGSNTNYADSVKCRF	FTFSNNAMNWVRQA-GAGLEWVSTLSY-DGSNTYYADSVKGRE	FTFSNYAMTWVRQAFGKGLEWVSVISS-VGSNTYYADSVKGRF	FTFSSYAMNWVRÇAFGKGLEWVSAILS-DGSSTSYADSVKGRF	FTFSSYAS-WVRÇAPGKGLEWVSGISG-DGSNTFYADSVKGFFTISRONSKNTLYLQMNSLRAEDTAVYYCARYDN	-FTFNSYAMIWVRÇA-GGGCLEWVSAIKS-DGSNTYYADSVKCRF=TSRONSKNTLYLQMNSLRAEDTAVYYCAR-	SGFTFSNYSMIWVRQAPGKGLEWVSGLSY-NGSNIYYADSVKGKE	FTFSNYY1SWVRÇADGKGLEWVSGISY NGSSTNYADSVKGRFTTSRNSKNTLYDWNSLRAEDTAVYYCARWW	FTFYKYAMHWVRÇA>GKGLEWVSGIQY-DGSYTYYADSVKGRFTTSRNSKNTLYLQMNSLRAEDTAVYYAAR	FTFSSYWMHWVRÇA=GKGLEWVSSISY-DSSNTYYADSVKGRFTISKNNSKNTLYLQMNSLRAEDTAVYYCAR	FTFSSYSMHWVRÇADGKGLEWVSGISY SSSFTYYADSVKGRFTTSRDNSKNTLYDWMSLRAEDTAVYYOARAL	SG—PTFSSYSMEWVRQA-CKGLEWVSGLSY-SSSFTYYADSVKERFTTSKNTLYLQMNSTRAEDTAVYYCARAL— sc—ptfstammingtdealckaettsc—meeptykadsvyrdeftetsch	LOT PHABILMAN
	SCAASGFT		SG	SG																			SCAASGFT	SCAASG FT	SG	SG	SC	SG	SG	SG	SG	SG	-	1	ł	SG	SC-	SG	SG	-	-	SG	SCAASGFT	CARAGGT - E
	'VQPGGSLRL	VQPGGSLRL:	VQPGGSLRL	VQPGGSLRL	VQPGGSLRL:	'VQPGGSLRL	'VXPSQTLSE'	VXPSQTLSE	VXZSQTLSZ	"LSLTUS-ZV"	'VXPSQTLSE'	'\Z>S\TT\S^;	TSTI.ÖS-YA	'VXPSQTLSE	LSTTQSEAU	'VXPSQTLSE'	'\Z>SQTI.S'.'	LSJTQS-YV	VX2SQTLSE	VXPSQTLST	ZXXPGASVIKV:	7442GESLRT	VQPGGSLRL:	VQPGGSLRL	VQPGGSLRL	VQPGGSLRL:	.VQPGGSI.R".	VQ2GGSLR_	NQPGGSLRL	VQPGGSLRL	'AQPGGSLRL'	.VQPGGST.R".	VQ2GGSLR.	'AQPGGSLRL'	VQPGGSLRL	'VQPGGSLRL	VQPGGSLRL	'VQ2GGSLR	'AQPGGSLRE'	VQPGGSLRL:	'VQPGGSLRL	'VQPGGSLRE'	VQPGGSLRI.	1 X X 1 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2
	QVQLVESGGGLVQPGGSLRLSCAA	QVQLVESGGGLVQPGGSLRLSCAA	QVQLVESGGGLVQPGGSLRLSCAA	QVQIVESGGGLVQPGGSLRLSCAA	QVQIVESGGGLVQPGGSLRLSCAA	2V21VESGGLVQPGGSLRISCAA	QVQLQQSGPGLVKPSQTLSLTCAL	QVQLQQSGFGLVXPSQTLSLTCAI	QVQLQQSGFGLVX7SQTLSTCAI	QVQLQQSGFGLVK-SQTLS_TCA1	QVQLQQSGFGLVXPSQTLSLTCAI	QVQ1 QQSGPG1,VX>SQT1,S_,TCAT	QVQLQQSGPGLVA2SQTLSLTCAL	QVQLQQSCPCLVKPSQTLSLTCAI	QVQLQQSGFGLVA2SQTLS_TCA1	QVQLQQSGFGLVXPSQTLSLTCAI	QVQI QQSGPGI,VX>SQTI,S_,TCAT	QVQLQQSGFGLVK-SQTLS_TCAL	QVQLQQSGFGLVX?SQTLSLTCAI	QVQLQQSGFGLVKPSQTLSLTCAI	QVQL VQSGAEVKKPGKA	QVQI VQSGAHVKKPGRSI,KTSCKG	QVQLVESGGGLVQPGGSLRLSCAA	QVQLVESGGGLVQPGGSLRLSCAA	QVQLVESGGGLVQPGGSLRLSC/A	QVQIVESGGGLVQPGGSLRLSCAA	QVQ1 VF SCCC1.VQ>CCS1.R7.SCAA.	QVQLVESGGGLVQZGGSLRLSCAA	QVQIVESGGGLVQPGGSLRLSCAA	QVQLVESGGGLVQPGGSLRLSCAA	QVQLVESGGGLVQPGGSLRLSCAA	QVQT VFSGGGT,VQ>GGST,R^,SCAA	QVQLVESGGGLVQZGGSLKLSCAASG	QVQIVESGGGLVQPGGSLRLSCAA	QVQIVESGGGLVQPGGSLRLSCAASG	QVQLVESGGGLVQPGGSLKLSCAA	QVQLVESCCCLVQPCCSLRLSCAA	QVQLVESGGGLVQ2GGSLR_SCAA.	QVQLVESGGGLVQPGGSLRLSCAA	QVQLVESGGGLVQPGGSLRLSCAA	QVQIVESGGGLVQPGGSLRLSCAASG-	QVQIVESGGGLVQPGGSLRLSCAA	QVQLVESGGGLVQPGGSLRLSCAA	ベマベエマドロロロロ
	TF-2A1C	TP-3B3	TP-2G4	P 2A5.1	TP-4A9	TP-2A8	TP-2B3	TP-2B9	TP 2H10	TP-3B4	TP-2C7	TP-2E3	TP-3C3	TP-2G5	T.P-4B7	TP-2G6	TP-3C2	TP-2D7	TP 3G1	TP-2E5	TP-2B8	TP-3F1	TP-3A3	TP 4E8	TP-4A7	TP-4H8	TP-2A6	TP-2C1	TP 3G3	TP-2B1	TP-2G7	TP-3H2	TP-2A2	TP-3E1	TP-2G2	TP-3D3	TP-2C9	TP-284	TP 3A2	TP-2F9	TP-3A4	TP 3C1	TP-3F2	055-47

Figure 8

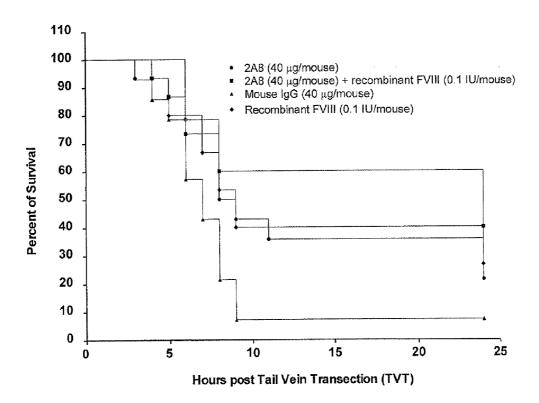
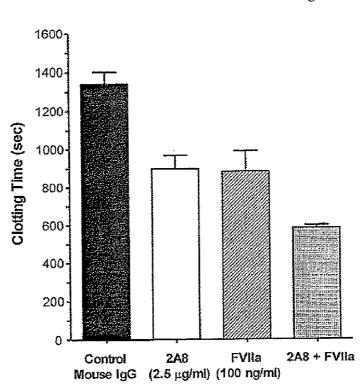


Figure 9



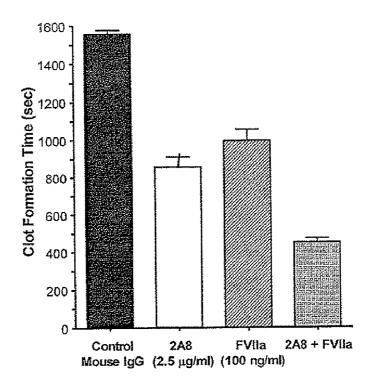
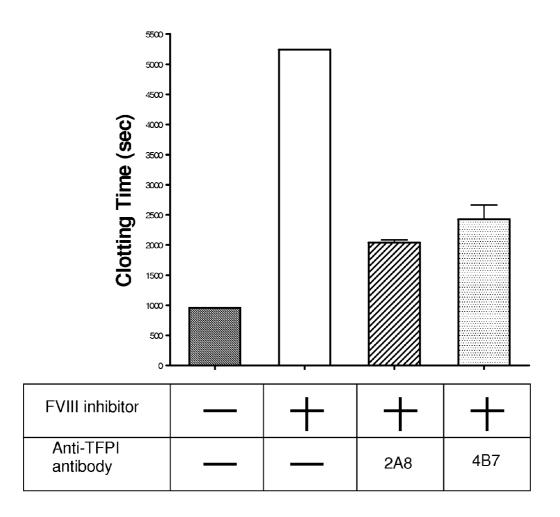


Figure 10



# MONOCLONAL ANTIBODIES AGAINST TISSUE FACTOR PATHWAY INHIBITOR

### SEQUENCE LISTING SUBMISSION

[0001] The Sequence Listing associated with this application is filed in electronic format via EFS-Web and hereby incorporated by reference into the specification in its entirety. The name of the text file containing the Sequence Listing is MSB7329PCT\_Sequence\_Listing\_ST25.

#### FIELD OF THE EMBODIMENTS

[0002] Provided are isolated monoclonal antibodies and fragments thereof that bind human tissue factor pathway inhibitor (TFPI) and related inventions.

# BACKGROUND [0003] Blood coagulation is a process by which blood

forms stable clots to stop bleeding. The process involves a number of proenzymes and procofactors (or "coagulation factors") that are circulating in the blood. Those proenzymes and procofactors interact through several pathways through which they are converted, either sequentially or simultaneously, to the activated form. Ultimately, the process results in the activation of prothrombin to thrombin by activated Factor X (FXa) in the presence of Factor Va, ionic calcium, and platelets. The activated thrombin in turn induces platelet aggregation and converts fibrinogen into fibrin, which is then cross linked by activated Factor XIII (FXIIIa) to form a clot. [0004] The process leading to the activation of Factor X can be carried out by two distinct pathways: the contact activation pathway (formerly known as the intrinsic pathway) and the tissue factor pathway (formerly known as the extrinsic pathway). It was previously thought that the coagulation cascade consisted of two pathways of equal importance joined to a common pathway. It is now known that the primary pathway for the initiation of blood coagulation is the tissue factor pathway.

[0005] Factor X can be activated by tissue factor (TF) in combination with activated Factor VII (FVIIa). The complex of Factor VIIa and its essential cofactor, TF, is a potent initiator of the clotting cascade.

[0006] The tissue factor pathway of coagulation is negatively controlled by tissue factor pathway inhibitor ("TFPI"). TFPI is a natural, FXa-dependent feedback inhibitor of the FVIIa/TF complex. It is a member of the multivalent Kunitz-type serine protease inhibitors. Physiologically, TFPI binds to activated Factor X (FXa) to form a heterodimeric complex, which subsequently interacts with the FVIIa/TF complex to inhibit its activity, thus shutting down the tissue factor pathway of coagulation. In principle, blocking TFPI activity can restore FXa and FVIIa/TF activity, thus prolonging the duration of action of the tissue factor pathway and amplifying the generation of FXa, which is the common defect in hemophilia A and B.

[0007] Indeed, some preliminary experimental evidence has indicated that blocking the TFPI activity by antibodies against TFPI normalizes the prolonged coagulation time or shortens the bleeding time. For instance, Nordfang et al. showed that the prolonged dilute prothrombin time of hemophilia plasma was normalized after treating the plasma with antibodies to TFPI (Thromb. Haemost., 1991, 66 (4): 464-467). Similarly, Erhardtsen et al. showed that the bleeding time in hemophilia A rabbit model was significantly short-

ened by anti-TFPI antibodies (Blood Coagulation and Fibrinolysis, 1995, 6: 388-394). These studies suggest that inhibition of TFPI by anti-TFPI antibodies may be useful for the treatment of hemophilia A or B. Only polyclonal anti-TFPI antibody was used in these studies.

[0008] Using hybridoma techniques, monoclonal antibodies against recombinant human TFPI (rhTFPI) were prepared and identified. See Yang et al., Chin. Med. J., 1998, 111 (8): 718-721. The effect of the monoclonal antibody on dilute prothrombin time (PT) and activated partial thromboplastin time (APTT) was tested. Experiments showed that anti-TFPI monoclonal antibody shortened dilute thromboplastin coagulation time of Factor IX deficient plasma. It is suggested that the tissue factor pathway plays an important role not only in physiological coagulation but also in hemorrhage of hemophilia (Yang et al., Hunan Yi Ke Da Xue Xue Bao, 1997, 22 (4): 297-300).

[0009] U.S. Pat. No. 7,015,194 to Kjalke et al. discloses compositions comprising FVIIa and a TFPI inhibitor, including polyclonal or monoclonal antibodies, or a fragment thereof, for treatment or prophylaxis of bleeding episodes or coagulative treatment. The use of such composition to reduce clotting time in normal mammalian plasma is also disclosed. It is further suggested that a Factor VIII or a variant thereof may be included in the disclosed composition of FVIIa and TFPI inhibitor. A combination of FVIII or Factor IX with TFPI monoclonal antibody is not suggested.

[0010] In addition to the treatment for hemophilia, it has also been suggested that TFPI inhibitors, including polyclonal or monoclonal antibodies, can be used for cancer treatment (see U.S. Pat. No. 5,902,582 to Hung).

[0011] Accordingly, antibodies specific for TFPI are needed for treating hematological diseases and cancer.

[0012] Generally, therapeutic antibodies for human diseases have been generated using genetic engineering to create murine, chimeric, humanized or fully human antibodies. Murine monoclonal antibodies were shown to have limited use as therapeutic agents because of a short serum half-life, an inability to trigger human effector functions, and the production of human antimouse-antibodies. Brekke and Sandlie, "Therapeutic Antibodies for Human Diseases at the Dawn of the Twenty-first Century," Nature 2, 53, 52-62 (January 2003). Chimeric antibodies have been shown to give rise to human anti-chimeric antibody responses. Humanized antibodies further minimize the mouse component of antibodies. However, a fully human antibody avoids the immunogenicity associated with murine elements completely. Thus, there is a need to develop fully human antibodies to avoid the immunogenicity associated with other forms of genetically engineered monoclonal antibodies. In particular, chronic prophylactic treatment such as would be required for hemophilia treatment with an anti-TFPI monoclonal antibody has a high risk of development of an immune response to the therapy if an antibody with a murine component or murine origin is used due to the frequent dosing required and the long duration of therapy. For example, antibody therapy for hemophilia A may require weekly dosing for the lifetime of a patient. This would be a continual challenge to the immune system. Thus, the need exists for a fully human antibody for antibody therapy for hemophilia and related genetic and acquired deficiencies or defects in coagulation.

[0013] Therapeutic antibodies have been made through hybridoma technology described by Koehler and Milstein in "Continuous Cultures of Fused Cells Secreting Antibody of

Predefined Specificity," *Nature* 256, 495-497 (1975). Fully human antibodies may also be made recombinantly in prokaryotes and eukaryotes. Recombinant production of an antibody in a host cell rather than hybridoma production is preferred for a therapeutic antibody. Recombinant production has the advantages of greater product consistency, likely higher production level, and a controlled manufacture that minimizes or eliminates the presence of animal-derived proteins. For these reasons, it is desirable to have a recombinantly produced monoclonal anti-TFPI antibody.

#### **SUMMARY**

[0014] Monoclonal antibodies to human tissue factor pathway inhibitor (TFPI) are provided. Further provided are the isolated nucleic acid molecules encoding the same. Pharmaceutical compositions comprising the anti-TFPI monoclonal antibodies and methods of treatment of genetic and acquired deficiencies or defects in coagulation such as hemophilia A and B are also provided. Also provided are methods for shortening the bleeding time by administering an anti-TFPI monoclonal antibody to a patient in need thereof. Methods for producing a monoclonal antibody that binds human TFPI according to the present invention are also provided.

#### BRIEF DESCRIPTION OF THE DRAWINGS

[0015] FIG. 1: The binding activity of representative examples of Fabs, selected from the panning and screening, to human TFPI ("h-TFPI") and mouse TFPI ("m-TFPI"). A control Fab against Estradiol-BSA ("EsB") and 12 Fabs (1-4 and 6-13) selected from panning TFPI were tested. Y-axis denotes fluorescence units of ELISA results.

[0016] FIG. 2: The dose-dependent in vitro functional activity of four representative anti-TFPI antibodies (4B7: TP-4B7, 2A8: TP-2A8, 2G6: TP-2G6, 2G7: TP-2G7) obtained from the panning and screening of a human antibody library as shown by their shortening dPT. The experiment involved 0.5 ug/mL of mTFPI spiked into TFPI depleted plasma.

[0017] FIG. 3: The in vitro functional activity of anti-TFPI Fab. Fab-2A8 (from TP-2A8), as tested in ROTEM assay.

[0018] FIG. 4: The binding activity to human TFPI and mouse TFPI of clones TP-2G6 ("2G6") after the conversion to IgG. A: IgG-2G6 binding to mouse TFPI; □: IgG-2G6 binding to human TFPI; ▲: control IgG binding to mouse TFPI; ■: control IgG binding to human IgG.

[0019] FIG. 5: The anti-TFPI antibodies TP-2A8 ("2A8"), TP-3G1 ("3G1"), and TP-3C2 ("3C2") shortened the whole blood clotting time in hemophilia A mice as tested in ROTEM assay. Each dot represents one individual hemophilia A mouse.

[0020] FIG. 6: The amino acid sequence alignment between the variable light chains of anti-TFPI monoclonal antibodies TP-2A10 (SEQ ID NO: 18), TP-2B1 (SEQ ID NO: 22), TP-2A2 (SEQ ID NO: 2), TP-2G2 (SEQ ID NO: 66), TP-2A5.1 (SEQ ID NO: 6), TP-3A3 (SEQ ID NO: 98), TP-2A8 (SEQ ID NO: 14), TP-2B8 (SEQ ID NO: 34), TP-2G7 (SEQ ID NO: 82), TP-4H8 (SEQ ID NO: 170), TP-2G4 (SEQ ID NO: 70), TP-3F2 (SEQ ID NO: 134), TP-2A6 (SEQ ID NO: 10), TP-3A2 (SEQ ID NO: 94), TP-2C1 (SEQ ID NO: 42), TP-3E1 (SEQ ID NO: 126), TP-3F1 (SEQ ID NO: 130), TP-3D3 (SEQ ID NO: 122), TP-4A7 (SEQ ID NO: 150), TP-4G8 (SEQ ID NO: 166), TP-2B3 (SEQ ID NO: 62),

TP-2G5 (SEQ ID NO: 74), TP-2G6 (SEQ ID NO: 78), TP-2H10 (SEQ ID NO: 90), TP-2B9 (SEQ ID NO: 38), TP-2C7 (SEQ ID NO: 46), TP-3G3 (SEQ ID NO: 142), TP-3C2 (SEQ ID NO: 114), TP-3B4 (SEQ ID NO: 110), TP-2E5 (SEQ ID NO: 58), TP-3C3 (SEQ ID NO: 118), TP-3G1 (SEQ ID NO: 138), TP-2D7 (SEQ ID NO: 50), TP-4B7 (SEQ ID NO: 158), TP-2E3 (SEQ ID NO: 54), TP-2G9 (SEQ ID NO: 86), TP-3C1 (SEQ ID NO: 86), TP-3A4 (SEQ ID NO: 102), TP-2B4 (SEQ ID NO: 30), TP-3H2 (SEQ ID NO: 146), TP-4A9 (SEQ ID NO: 154), TP-4E8 (SEQ ID NO: 162), and TP-3B3 (SEQ ID NO: 106). [0021] FIG. 7: The amino acid sequence alignment between the variable heavy chains of anti-TFPI monoclonal antibodies TP-2A10 (SEQ ID NO: 20), TP-3B3 (SEQ ID NO: 108), TP-2G4 (SEQ ID NO: 72), TP-2A5.1 (SEQ ID NO: 8), TP-4A9 (SEQ ID NO: 156), TP-2A8 (SEQ ID NO: 16), TP-2B3 (SEQ ID NO: 28), TP-2B9 (SEQ ID NO: 40), TP-2H10 (SEQ ID NO: 92), TP-3B4 (SEQ ID NO: 112), TP-2C7 (SEQ ID NO: 48), TP-2E3 (SEQ ID NO: 56), TP-3C3 (SEQ ID NO: 120), TP-2G5 (SEQ ID NO: 76), TP-4B7 (SEQ ID NO: 160), TP-2G6 (SEQ ID NO: 80), TP-3C2 (SEQ ID NO: 116), TP-2D7 (SEQ ID NO: 52), TP-3G1 (SEQ ID NO: 140), TP-2E5 (SEQ ID NO: 60), TP-2B8 (SEQ ID NO: 36), TP-3F1 (SEQ ID NO: 132), TP-3A3 (SEQ ID NO: 100), TP-4E8 (SEQ ID NO: 164), TP-4A7 (SEQ ID NO: 152), TP-4H8 (SEQ ID NO: 172), TP-2A6 (SEQ ID NO: 12), TP-2C1 (SEQ ID NO: 44), TP-3G3 (SEQ ID NO: 144), TP-2B1 (SEQ ID NO: 24), TP-2G7 (SEQ ID NO: 84), TP-3H2 (SEQ ID NO: 148), TP-2A2 (SEQ ID NO: 4), TP-3E1 (SEQ ID NO: 128), TP-2G2 (SEQ ID NO: 68), TP-3D3 (SEQ ID NO: 124), TP-2G9 (SEQ ID NO: 88), TP-2B4 (SEQ ID NO: 32), TP-3A2 (SEQ ID NO: 96), TP-2F9 (SEQ ID NO: 64), TP-3A4 (SEQ ID NO: 104), TP-3C1 (SEQ ID NO: 136), TP-3F2 (SEQ ID NO: 136), and TP-4G8 (SEQ ID NO: 168). [0022] FIG. 8: Graph showing the survival rate over 24 hours post-tail vein transection for mice treated with (1) the anti-TFPI antibody TP-2A8 ("2A8"), (2) 2A8 and recombinant factor VIII, (3) mouse IgG, and (4) recombinant factor

[0023] FIG. 9: Graphs showing clotting time and clot formation time assays for mice treated with the anti-TFPI antibody TP-2A8 ("2A8"), factor VIIa, and the combination of 2A8 and factor VIIa.

[0024] FIG. 10: Graph showing clotting time for normal human blood treated with a FVIII inhibitor with the anti-TFPI antibody TP-2A8 ("2A8")and anti-TFPI antibody TP-4B7 ("4B7") as compared to FVIII inhibitor alone.

#### DETAILED DESCRIPTION

#### Definitions

[0025] The term "tissue factor pathway inhibitor" or "TFPI" as used herein refers to any variant, isoform and species homolog of human TFPI that is naturally expressed by cells. In a preferred embodiment of the invention, the binding of an antibody of the invention to TFPI reduces the blood clotting time.

[0026] As used herein, an "antibody" refers to a whole antibody and any antigen binding fragment (i.e., "antigenbinding portion") or single chain thereof. The term includes a full-length immunoglobulin molecule (e.g., an IgG antibody) that is naturally occurring or formed by normal immunoglobulin gene fragment recombinatorial processes, or an immu-

nologically active portion of an immunoglobulin molecule, such as an antibody fragment, that retains the specific binding activity. Regardless of structure, an antibody fragment binds with the same antigen that is recognized by the full-length antibody. For example, an anti-TFPI monoclonal antibody fragment binds to an epitope of TFPI. The antigen-binding function of an antibody can be performed by fragments of a full-length antibody. Examples of binding fragments encompassed within the term "antigen-binding portion" of an antibody include (i) a Fab fragment, a monovalent fragment consisting of the  $V_L, V_H, C_L$  and  $C_{H1}$  domains; (ii) a  $F(ab')_2$ fragment, a bivalent fragment comprising two Fab fragments linked by a disulfide bridge at the hinge region; (iii) a Fd fragment consisting of the  $V_H$  and  $C_{H1}$  domains; (iv) a Fv fragment consisting of the  $V_L$  and  $V_H$  domains of a single arm of an antibody, (v) a dAb fragment (Ward et al., (1989) Nature 341:544-546), which consists of a  $V_H$  domain; and (vi) an isolated complementarity determining region (CDR). Furthermore, although the two domains of the Fv fragment,  $V_L$ and  $V_H$ , are coded for by separate genes, they can be joined, using recombinant methods, by a synthetic linker that enables them to be made as a single protein chain in which the  $V_L$  and  $V_H$  regions pair to form monovalent molecules (known as single chain Fv (scFv); see e.g., Bird et al. (1988) Science 242:423-426; and Huston et al (1988) Proc. Natl. Acad. Sci. USA 85:5879-5883). Such single chain antibodies are also intended to be encompassed within the term "antigen-binding portion" of an antibody. These antibody fragments are obtained using conventional techniques known to those with skill in the art, and the fragments are screened for utility in the same manner as are intact antibodies.

[0027] As used herein, the terms "inhibits binding" and "blocks binding" (e.g., referring to inhibition/blocking of binding of TFPI ligand to TFPI) are used interchangeably and encompass both partial and complete inhibition or blocking. Inhibition and blocking are also intended to include any measurable decrease in the binding affinity of TFPI to a physiological substrate when in contact with an anti-TFPI antibody as compared to TFPI not in contact with an anti-TFPI antibody, e.g., the blocking of the interaction of TFPI with factor Xa or blocking the interaction of a TFPI-factor Xa complex with tissue factor, factor VIIa or the complex of tissue factor/factor VIIa by at least about 10%, 20%, 30%, 40%, 50%, 60%, 70%, 80%, 90%, 95%, 96%, 97%, 98%, 99%, or 100%.

[0028] The terms "monoclonal antibody" or "monoclonal antibody composition" as used herein refer to a preparation of antibody molecules of single molecular composition. A monoclonal antibody composition displays a single binding specificity and affinity for a particular epitope. Accordingly, the term "human monoclonal antibody" refers to antibodies displaying a single binding specificity which have variable and constant regions derived from human germline immunoglobulin sequences. The human antibodies of the invention may include amino acid residues not encoded by human germline immunoglobulin sequences (e.g., mutations introduced by random or site-specific mutagenesis in vitro or by somatic mutation in vivo).

[0029] An "isolated antibody," as used herein, is intended to refer to an antibody which is substantially free of other antibodies having different antigenic specificities (e.g., an isolated antibody that binds to TFPI is substantially free of antibodies that bind antigens other than TFPI). An isolated antibody that binds to an epitope, isoform or variant of human TFPI may, however, have cross-reactivity to other related

antigens, e.g., from other species (e.g., TFPI species homologs). Moreover, an isolated antibody may be substantially free of other cellular material and/or chemicals.

[0030] As used herein, "specific binding" refers to antibody binding to a predetermined antigen. Typically, the antibody binds with an affinity of at least about 10<sup>5</sup> M<sup>-1</sup> and binds to the predetermined antigen with an affinity that is higher, for example at least two-fold greater, than its affinity for binding to an irrelevant antigen (e.g., BSA, casein) other than the predetermined antigen or a closely-related antigen. The phrases "an antibody recognizing an antigen" and "an antibody specific for an antigen" are used interchangeably herein with the term "an antibody which binds specifically to an antigen."

**[0031]** As used herein, the term "high affinity" for an IgG antibody refers to a binding affinity of at least about  $10^7 M^{-1}$ , in some embodiments at least about  $10^8 M^{-1}$ , in some embodiments at least about  $10^9 M^{-1}$ ,  $10^{10} M^{-1}$ ,  $10^{11} M^{-1}$  or greater, e.g., up to  $10^{13} M^{-1}$  or greater.

[0032] However, "high affinity" binding can vary for other antibody isotypes. For example, "high affinity" binding for an IgM isotype refers to a binding affinity of at least about  $1.0\times10^7 M^{-1}$ . As used herein, "isotype" refers to the antibody class (e.g., IgM or IgG1) that is encoded by heavy chain constant region genes.

[0033] "Complementarity-determining region" or "CDR" refers to one of three hypervariable regions within the variable region of the heavy chain or the variable region of the light chain of an antibody molecule that form the N-terminal antigen-binding surface that is complementary to the three-dimensional structure of the bound antigen. Proceeding from the N-terminus of a heavy or light chain, these complementarity-determining regions are denoted as "CDR1," "CDR2," and "CDR3," respectively. CDRs are involved in antigenantibody binding, and the CDR3 comprises a unique region specific for antigen-antibody binding. An antigen-binding site, therefore, may include six CDRs, comprising the CDR regions from each of a heavy and a light chain V region.

[0034] As used herein, "conservative substitutions" refers to modifications of a polypeptide that involve the substitution of one or more amino acids for amino acids having similar biochemical properties that do not result in loss of a biological or biochemical function of the polypeptide. A "conservative amino acid substitution" is one in which the amino acid residue is replaced with an amino acid residue having a similar side chain. Families of amino acid residues having similar side chains have been defined in the art. These families include amino acids with basic side chains (e.g., lysine, arginine, histidine), acidic side chains (e.g., aspartic acid, glutamic acid), uncharged polar side chains (e.g., glycine, asparagine, glutamine, serine, threonine, tyrosine, cysteine), nonpolar side chains (e.g., alanine, valine, leucine, isoleucine, proline, phenylalanine, methionine, tryptophan), betabranched side chains (e.g., threonine, valine, isoleucine), and aromatic side chains (e.g., tyrosine, phenylalanine, tryptophan, histidine). It is envisioned that the antibodies of the present invention may have conservative amino acid substitutions and still retain activity.

[0035] For nucleic acids and polypeptides, the term "substantial homology" indicates that two nucleic acids or two polypeptides, or designated sequences thereof, when optimally aligned and compared, are identical, with appropriate nucleotide or amino acid insertions or deletions, in at least about 80% of the nucleotides or amino acids, usually at least

about 85%, preferably about 90%, 91%, 92%, 93%, 94%, or 95%, more preferably at least about 96%, 97%, 98%, 99%, 99.1%, 99.2%, 99.3%, 99.4%, or 99.5% of the nucleotides or amino acids. Alternatively, substantial homology for nucleic acids exists when the segments will hybridize under selective hybridization conditions to the complement of the strand. The invention includes nucleic acid sequences and polypeptide sequences having substantial homology to the specific nucleic acid sequences and amino acid sequences recited herein.

[0036] The percent identity between two sequences is a function of the number of identical positions shared by the sequences (i.e., % homology=# of identical positions/total # of positions×100), taking into account the number of gaps, and the length of each gap, which need to be introduced for optimal alignment of the two sequences. The comparison of sequences and determination of percent identity between two sequences can be accomplished using a mathematical algorithm, such as without limitation the AlignXTM module of VectorNTITM (Invitrogen Corp., Carlsbad, Calif.). For AlignX<sup>TM</sup>, the default parameters of multiple alignment are: gap opening penalty: 10; gap extension penalty: 0.05; gap separation penalty range: 8; % identity for alignment delay: 40. (further details found at http://www.invitrogen.com/site/ us/en/home/LINNEA-Online-Guides/LINNEA-Communities/Vector-NTI-Community/Sequence-analysis-and-datamanagement-software-for-PCs/AlignX-Module-for-Vector-NTI-Advance.reg.us.html).

[0037] Another method for determining the best overall match between a query sequence (a sequence of the present invention) and a subject sequence, also referred to as a global sequence alignment, can be determined using the CLUST-ALW computer program (Thompson et al., Nucleic Acids

Research, 1994, 2 (22): 4673-4680), which is based on the algorithm of Higgins et al., (Computer Applications in the Biosciences (CABIOS), 1992, 8 (2): 189-191). In a sequence alignment the query and subject sequences are both DNA sequences. The result of said global sequence alignment is in percent identity. Preferred parameters used in a CLUSTALW alignment of DNA sequences to calculate percent identity via pairwise alignments are: Matrix=IUB, k-tuple=1, Number of Top Diagonals=5, Gap Penalty=3, Gap Open Penalty=10, Gap Extension Penalty=0.1. For multiple alignments, the following CLUSTALW parameters are preferred: Gap Opening Penalty=10, Gap Extension Parameter=0.05; Gap Separation Penalty Range=8; % Identity for Alignment Delay=40.

[0038] The nucleic acids may be present in whole cells, in a cell lysate, or in a partially purified or substantially pure form. A nucleic acid is "isolated" or "rendered substantially pure" when purified away from other cellular components with which it is normally associated in the natural environment. To isolate a nucleic acid, standard techniques such as the following may be used: alkaline/SDS treatment, CsC1 banding, column chromatography, agarose gel electrophoresis and others well known in the art.

#### Monoclonal Antibodies

[0039] Forty-four TFPI-binding antibodies were identified from panning and screening of human antibody libraries against human TFPI. The heavy chain variable region and light chain variable region of each monoclonal antibody were sequenced and their CDR regions were identified. The sequence identifier numbers ("SEQ ID NO") correspond to these regions of each monoclonal antibody are summarized in Table 1.

TABLE 1

Summary of the sequence identifier numbers ("SEQ ID NO") of the heavy chain variable region ("VH") and light chain variable region ("VL") of each TFPI-binding monoclonal antibodies. The sequence identifier numbers for the CDR regions ("CDR1," "CDR2," and "CDR3") of each heavy and light chain are also provided.

	V	L	V	H		VL			VH	
Clone	N.A.	A.A.	N.A.	A.A.	CDR1	CDR2	CDR3	CDR1	CDR2	CDR3
TP-2A2	1	2	3	4	173	216	259	302	345	388
TP-2A5.1	5	6	7	8	174	217	260	303	346	389
TP-2A6	9	10	11	12	175	218	261	304	347	390
TP-2A8	13	14	15	16	176	219	262	305	348	391
TP-2A10	17	18	19	20	177	220	263	306	349	392
TP-2B1	21	22	23	24	178	221	264	307	350	393
TP-2B3	25	26	27	28	179	222	265	308	351	394
TP-2B4	29	30	31	32	180	223	266	309	352	395
TP-2B8	33	34	35	36	181	224	267	310	353	396
TP-2B9	37	38	39	40	182	225	268	311	354	397
TP-2C1	41	42	43	44	183	226	269	312	355	398
TP-2C7	45	46	47	48	184	227	270	313	356	399
TP-2D7	49	50	51	52	185	228	271	314	357	400
TP-2E3	53	54	55	56	186	229	272	315	358	401
TP-2E5	57	58	59	60	187	230	273	316	359	402
TP-2F9	61	62	63	64	188	231	274	317	360	403
TP-2G2	65	66	67	68	189	232	275	318	361	404
TP-2G4	69	70	71	72	190	233	276	319	362	405
TP-2G5	73	74	75	76	191	234	277	320	363	406
TP-2G6	77	78	79	80	192	235	278	321	364	407
TP-2G7	81	82	83	84	193	236	279	322	365	408
TP-2G9	85	86	87	88	194	237	280	323	366	409
TP-2H10	89	90	91	92	195	238	281	324	367	410
TP-3A2	93	94	95	96	196	239	282	325	368	411

TABLE 1-continued

Summary of the sequence identifier numbers ("SEQ ID NO") of the heavy chain variable region ("VH") and light chain variable region ("VL") of each TFPI-binding monoclonal antibodies. The sequence identifier numbers for the CDR regions ("CDR1," "CDR2," and "CDR3") of each heavy and light chain are also provided.

	V	<u>L</u>	V	Η		VL			VH	
Clone	N.A.	A.A.	N.A.	A.A.	CDR1	CDR2	CDR3	CDR1	CDR2	CDR3
TP-3A3	97	98	99	100	197	240	283	326	369	412
TP-3A4	101	102	103	104	198	241	284	327	370	413
TP-3B3	105	106	107	108	199	242	285	328	371	414
TP-3B4	109	110	111	112	200	243	286	329	372	415
TP-3C2	113	114	115	116	201	244	287	330	373	416
TP-3C3	117	118	119	120	202	245	288	331	374	417
TP-3D3	121	122	123	124	203	246	289	332	375	418
TP-3E1	125	126	127	128	204	247	290	333	376	419
TP-3F1	129	130	131	132	205	248	291	334	377	420
TP-3F2	133	134	135	136	206	249	292	335	378	421
TP-3G1	137	138	139	140	207	250	293	336	379	422
TP-3G3	141	142	143	144	208	251	294	337	380	423
TP-3H2	145	146	147	148	209	252	295	338	381	424
TP-4A7	149	150	151	152	210	253	296	339	382	425
TP-4A9	153	154	155	156	211	254	297	340	383	426
TP-4B7	157	158	159	160	212	255	298	341	384	427
TP-4E8	161	162	163	164	213	256	299	342	385	428
TP-4G8	165	166	167	168	214	257	300	343	386	429
TP-4H8	169	170	171	172	215	258	301	344	387	430
TP-3C1	85	86	135	136	194	237	280	335	378	421

N.A.: nucleic acid sequence; A.A.: amino acid sequence.

[0040] In one embodiment, provided is an isolated monoclonal antibody that binds to human tissue factor pathway inhibitor, wherein the antibody comprises a CDR3 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 388-430. These CDR3s are identified from the heavy chains of the antibodies identified during panning and screening. In a further embodiment, this antibody further comprises (a) a CDR1 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 302-344, (b) a CDR2 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 345-387, or (c) both a CDR1 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 302-344 and a CDR2 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 345-387.

[0041] In another embodiment, provided are antibodies that share a CDR3 from one of the light chains of the antibodies identified during panning and screening. Thus, the present invention is directed to an isolated monoclonal antibody that binds to human tissue factor pathway inhibitor, wherein the antibody comprises a CDR3 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 259-301. In further embodiments, the antibody further comprises (a) a CDR1 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 173-215, (b) a CDR2 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 216-258, or (c) both a CDR1 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 173-215 and a CDR2 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 216-258.

[0042] In another embodiment, the antibody comprises a CDR3 from a heavy chain and a CDR3 from a light chain of the antibodies identified from screening and panning. Thus,

provided is an antibody that binds to human tissue factor pathway inhibitor, wherein the antibody comprises a CDR3 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 388-430 and a CDR3 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 259-301. In a further embodiment, the antibody further comprises (a) a CDR1 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 302-344, (b) a CDR2 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 345-387, (c) a CDR1 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 173-215, and/or (d) a CDR2 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 216-258.

[0043] In other specific embodiments, the antibody comprises heavy and light chain variable regions comprising:

[0044] (a) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 173, 216 and 259 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 302, 345 and 388;

[0045] (b) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 174, 217 and 260 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 303, 346 and 389;

[0046] (c) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 175, 218 and 261 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 304, 347 and 390;

[0047] (d) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 176, 219

- and 262 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 305, 348 and 391;
- [0048] (e) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 177, 220 and 263 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 306, 349 and 392:
- [0049] (f) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 178, 221 and 264 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 307, 350 and 393;
- [0050] (g) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 179, 222 and 265 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 308, 351 and 394:
- [0051] (h) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 180, 223 and 266 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 309, 352 and 395;
- [0052] (i) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 181, 224 and 267 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 310, 353 and 396;
- [0053] (j) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 182, 225 and 268 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 311, 354 and 397;
- [0054] (k) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 183, 226 and 269 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 312, 355 and 398;
- [0055] (1) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 184, 227 and 270 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 313, 356 and 399;
- [0056] (m) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 185, 228 and 271 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 314, 357 and 400;
- [0057] (n) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 186, 229 and 272 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 315, 358 and 401;
- [0058] (o) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 187, 230 and 273 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 316, 359 and 402;
- [0059] (p) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 188, 231 and 274 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 317, 360 and 403;
- [0060] (q) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 189, 232 and 275 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 318, 361 and 404;

- [0061] (r) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 190, 233 and 276 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 319, 362 and 405;
- [0062] (s) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 191, 234 and 277 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 320, 363 and 406:
- [0063] (t) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 192, 235 and 278 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 321, 364 and 407;
- [0064] (u) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 193, 236 and 279 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 322, 365 and 408:
- [0065] (v) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 194, 237 and 280 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 323, 366 and 409;
- [0066] (w) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 195, 238 and 281 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 324, 367 and 410;
- [0067] (x) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 196, 239 and 282 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 325, 368 and 411:
- [0068] (y) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 197, 240 and 283 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 326, 369 and 412;
- [0069] (z) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 198, 241 and 284 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 327, 370 and 413:
- [0070] (aa) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 199, 242 and 285 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 328, 371 and 414;
- [0071] (bb) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 200, 243 and 286 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 329, 372 and 415;
- [0072] (cc) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 201, 244 and 287 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 330, 373 and;
- [0073] (dd) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 202, 245 and 288 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 331, 374 and 417;

- [0074] (ee) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 203, 246 and 289 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 332, 375 and 418;
- [0075] (ff) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 204, 247 and 290 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 333, 376 and 419:
- [0076] (gg) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 205, 248 and 291 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 334, 377 and 420;
- [0077] (hh) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 206, 249 and 292 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 335, 378 and 421;
- [0078] (ii) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 207, 250 and 293 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 336, 379 and 422;
- [0079] (jj) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 208, 251 and 294 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 337, 380 and 423;
- [0080] (kk) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 209, 252 and 295 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 338, 381 and 424;
- [0081] (II) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 210, 253 and 296 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 339, 382 and 425;
- [0082] (mm) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 211, 254 and 297 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 340, 383 and 426;
- [0083] (nn) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 212, 255 and 298 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 341, 384 and 427;
- [0084] (00) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 213, 256 and 299 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 342, 385 and 428;
- [0085] (pp) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 214, 257 and 300 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 343, 386 and 429;
- [0086] (qq) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 215, 258

- and 301 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 344, 387 and 430; or
- [0087] (rr) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 194, 237 and 280 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 335, 378 and 421.
- [0088] In another embodiment, the invention is directed to antibodies comprising:
  - [0089] (a) a light chain variable region having the polypeptide sequence of SEQ ID NO: 2 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 4;
  - [0090] (b) a light chain variable region having the polypeptide sequence of SEQ ID NO: 6 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 8;
  - [0091] (c) a light chain variable region having the polypeptide sequence of SEQ ID NO: 10 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 12;
  - [0092] (d) a light chain variable region having the polypeptide sequence of SEQ ID NO: 14 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 16;
  - [0093] (e) a light chain variable region having the polypeptide sequence of SEQ ID NO: 18 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 20;
  - [0094] (f) a light chain variable region having the polypeptide sequence of SEQ ID NO: 22 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 24;
  - [0095] (g) a light chain variable region having the polypeptide sequence of SEQ ID NO: 26 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 28;
  - [0096] (h) a light chain variable region having the polypeptide sequence of SEQ ID NO: 30 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 32;
  - [0097] (i) a light chain variable region having the polypeptide sequence of SEQ ID NO: 34 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 36;
  - [0098] (j) a light chain variable region having the polypeptide sequence of SEQ ID NO: 38 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 40;
  - [0099] (k) a light chain variable region having the polypeptide sequence of SEQ ID NO: 42 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 44;
  - [0100] (l) a light chain variable region having the polypeptide sequence of SEQ ID NO: 46 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 48;
  - [0101] (m) a light chain variable region having the polypeptide sequence of SEQ ID NO: 50 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 52;

- [0102] (n) a light chain variable region having the polypeptide sequence of SEQ ID NO: 54 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 56;
- [0103] (o) a light chain variable region having the polypeptide sequence of SEQ ID NO: 58 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 60;
- [0104] (p) a light chain variable region having the polypeptide sequence of SEQ ID NO: 62 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 64;
- [0105] (q) a light chain variable region having the polypeptide sequence of SEQ ID NO: 66 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 68;
- [0106] (r) a light chain variable region having the polypeptide sequence of SEQ ID NO: 70 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 72;
- [0107] (s) a light chain variable region having the polypeptide sequence of SEQ ID NO: 74 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 76;
- [0108] (t) a light chain variable region having the polypeptide sequence of SEQ ID NO: 78 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 80;
- [0109] (u) a light chain variable region having the polypeptide sequence of SEQ ID NO: 82 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 84;
- [0110] (v) a light chain variable region having the polypeptide sequence of SEQ ID NO: 86 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 88;
- [0111] (w) a light chain variable region having the polypeptide sequence of SEQ ID NO: 90 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 92;
- [0112] (x) a light chain variable region having the polypeptide sequence of SEQ ID NO: 94 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 96;
- [0113] (y) a light chain variable region having the polypeptide sequence of SEQ ID NO: 98 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 100;
- [0114] (z) a light chain variable region having the polypeptide sequence of SEQ ID NO: 102 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 104;
- [0115] (aa) a light chain variable region having the polypeptide sequence of SEQ ID NO: 106 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 108;
- [0116] (bb) a light chain variable region having the polypeptide sequence of SEQ ID NO: 110 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 112;
- [0117] (cc) a light chain variable region having the polypeptide sequence of SEQ ID NO: 114 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 116;

- [0118] (dd) a light chain variable region having the polypeptide sequence of SEQ ID NO: 118 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 120;
- [0119] (ee) a light chain variable region having the polypeptide sequence of SEQ ID NO: 122 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 124;
- [0120] (ff) a light chain variable region having the polypeptide sequence of SEQ ID NO: 126 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 128;
- [0121] (gg) a light chain variable region having the polypeptide sequence of SEQ ID NO: 130 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 132;
- [0122] (hh) a light chain variable region having the polypeptide sequence of SEQ ID NO: 134 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 136;
- [0123] (ii) a light chain variable region having the polypeptide sequence of SEQ ID NO: 138 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 140;
- [0124] (jj) a light chain variable region having the polypeptide sequence of SEQ ID NO: 142 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 144;
- [0125] (kk) a light chain variable region having the polypeptide sequence of SEQ ID NO: 146 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 148;
- [0126] (II) a light chain variable region having the polypeptide sequence of SEQ ID NO: 150 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 152;
- [0127] (mm) a light chain variable region having the polypeptide sequence of SEQ ID NO: 154 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 156;
- [0128] (nn) a light chain variable region having the polypeptide sequence of SEQ ID NO: 158 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 160;
- [0129] (oo) a light chain variable region having the polypeptide sequence of SEQ ID NO: 162 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 164;
- [0130] (pp) a light chain variable region having the polypeptide sequence of SEQ ID NO: 166 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 168;
- [0131] (qq) a light chain variable region having the polypeptide sequence of SEQ ID NO: 170 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 172; or
- [0132] (rr) a light chain variable region having the polypeptide sequence of SEQ ID NO: 86 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 136.
- [0133] Also provided is an isolated monoclonal antibody that binds to human tissue factor pathway inhibitor, wherein the antibody comprises a human heavy chain variable region comprising an amino acid sequence having at least 89%,

90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 99.5% identity to an amino acid sequence selected from the group consisting of the amino acid sequences set forth in SEQ ID NO:4, SEQ ID NO:8, SEQ ID NO:12, SEQ ID NO:16, SEQ ID NO:20, SEQ ID NO:24, SEQ ID NO:28, SEQ ID NO:32, SEQ ID NO:36, SEQ ID NO:40, SEQ ID NO:44. SEQ ID NO:48, SEQ ID NO:52, SEQ ID NO:56, SEQ ID NO:60, SEQ ID NO:64, SEQ ID NO:68, SEQ ID NO:72, SEQ ID NO:76, SEQ ID NO:80, SEQ ID NO:84, SEQ ID NO:88, SEQ ID NO:92, SEQ ID NO:96, SEQ ID NO:100, SEQ ID NO:104, SEQ ID NO:108, SEQ ID NO:112, SEQ ID NO:116, SEQ ID NO:120, SEQ ID NO:124, SEQ ID NO:128, SEQ ID NO:132, SEQ ID NO:136, SEQ ID NO:140, SEQ ID NO:144, SEQ ID NO:148, SEQ ID NO:152, SEQ ID NO:156, SEQ ID NO:160, SEQ ID NO:164, SEQ ID NO:168, and SEQ ID NO:172.

[0134] Also provided is an isolated monoclonal antibody that binds to human tissue factor pathway inhibitor, wherein the antibody comprises a human light chain variable region comprising an amino acid sequence having at least 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 99.5% identity to an amino acid sequence selected from the group consisting of the amino acid sequences set forth in SEQ ID NO:2, SEQ ID NO:6, SEQ ID NO:10, SEQ ID NO:14, SEQ ID NO:18, SEQ ID NO:22, SEQ ID NO:26, SEQ ID NO:30, SEQ ID NO:34, SEQ ID NO:38, SEQ ID NO:42, SEQ ID NO:46, SEQ ID NO:50, SEQ ID NO:54, SEQ ID NO:58, SEQ ID NO:62, SEO ID NO:66, SEO ID NO:70, SEO ID NO:74, SEO ID NO:78, SEQ ID NO:82, SEQ ID NO:86, SEQ ID NO:90, SEQ ID NO:94, SEQ ID NO:98, SEQ ID NO:102, SEQ ID NO:106, SEQ ID NO:110, SEQ ID NO:114, SEQ ID NO:118, SEQ ID NO:122, SEQ ID NO:126, SEQ ID NO:130, SEQ ID NO:134, SEQ ID NO:138, SEQ ID NO:142, SEQ ID NO:146, SEQ ID NO:150, SEQ ID NO:154, SEQ ID NO:158, SEQ ID NO:162, SEQ ID NO:166, and SEQ ID NO:170.

[0135] In addition to relying on the antibody descriptions using the sequence identifiers discussed above, some embodiments may also be described by reference to the Fab clones isolated in the experiments described herein. In some embodiments, the recombinant antibodies comprise the heavy and/or light chain CDR3s of the following clones: TP-2A2, TP-2A5.1, TP-2A6, TP-2A8, TP-2A10, TP-2B1, TP-2B3, TP-2B4, TP-2B8, TP-2B9, TP-2C1, TP-2C7, TP-2D7, TP-2E3, TP-2E5, TP-2F9, TP-2G2, TP-2G4, TP-2G5, TP-2G6, TP-2G7, TP-2G9, TP-2H10, TP-3A2, TP-3A3, TP-3A4, TP-3B3, TP-3B4, TP-3C1, TP-3C2, TP-3C3, TP-3D3, TP-3E1, TP-3F1, TP-3F2, TP-3G1, TP-3G3, TP-3H2, TP-4A7, TP-4A9, TP-4B7, TP-4E8, TP-4G8, or TP-4H8. In some embodiments, the antibodies further can comprise the CDR2s of these antibodies and still further comprise the CDR1s of these antibodies. In other embodiments, the antibodies can further comprise any combinations of the CDRs.

[0136] Accordingly, in another embodiment, provided are anti-TFPI antibodies comprising: (1) human heavy chain framework regions, a human heavy chain CDR1 region, a human heavy chain CDR3 region, and a human heavy chain CDR3 region is the heavy chain CDR3 of TP-2A2, TP-2A5.1, TP-2A6, TP-2A8, TP-2A10, TP-2B1, TP-2B3, TP-2B4, TP-2B8, TP-2B9, TP-2C1, TP-2C7, TP-2D7, TP-2E3, TP-2E5, TP-2F9, TP-2G2, TP-2G4, TP-2G5, TP-2G6, TP-2G7, TP-2G9, TP-2H10, TP-3A2, TP-3A3, TP-3A4, TP-3B3,

TP-3B4, TP-3C1, TP-3C2, TP-3C3, TP-3D3, TP-3E1, TP-3F1, TP-3F2, TP-3G1, TP-3G3, TP-3H2, TP-4A7, TP-4A9, TP-4B7, TP-4E8, TP-4G8, or TP-4H8; and (2) human light chain framework regions, a human light chain CDR1 region, a human light chain CDR2 region, and a human light chain CDR3 region, wherein the human light chain CDR3 region is the light chain CDR3 of TP-2A2, TP-2A5.1, TP-2A6, TP-2A8, TP-2A10, TP-2B1, TP-2B3, TP-2B4, TP-2B8, TP-2B9, TP-2C1, TP-2C7, TP-2D7, TP-2E3, TP-2E5, TP-2F9, TP-2G2, TP-2G4, TP-2G5, TP-2G6, TP-2G7, TP-2G9, TP-2H10, TP-3A2, TP-3A3, TP-3A4, TP-3B3, TP-3B4, TP-3C1, TP-3C2, TP-3C3, TP-3D3, TP-3E1, TP-3F1, TP-3F2, TP-3G1, TP-3G3, TP-3H2, TP-4A7, TP-4A9, TP-4B7, TP-4E8, TP-4G8, or TP-4H8, wherein the antibody binds TFPI. The antibody may further comprise the heavy chain CDR2 and/or the light chain CDR2 of TP-2A2, TP-2A5.1, TP-2A6, TP-2A8, TP-2A10, TP-2B1, TP-2B3, TP-2B4, TP-2B8, TP-2B9, TP-2C1, TP-2C7, TP-2D7, TP-2E3, TP-2E5, TP-2F9, TP-2G2, TP-2G4, TP-2G5, TP-2G6, TP-2G7, TP-2G9, TP-2H10, TP-3A2, TP-3A3, TP-3A4, TP-3B3, TP-3B4, TP-3C1, TP-3C2, TP-3C3, TP-3D3, TP-3E1, TP-3F1, TP-3F2, TP-3G1, TP-3G3, TP-3H2, TP-4A7, TP-4A9, TP-4B7, TP-4E8, TP-4G8, or TP-4H8. The antibody may further comprise the heavy chain CDR1 and/or the light chain CDR1 of TP-2A2, TP-2A5.1, TP-2A6, TP-2A8, TP-2A10, TP-2B1, TP-2B3, TP-2B4, TP-2B8, TP-2B9, TP-2C1, TP-2C7, TP-2D7, TP-2E3, TP-2E5, TP-2F9, TP-2G2, TP-2G4, TP-2G5, TP-2G6, TP-2G7, TP-2G9, TP-2H10, TP-3A2, TP-3A3, TP-3A4, TP-3B3, TP-3B4, TP-3C1, TP-3C2, TP-3C3, TP-3D3, TP-3E1, TP-3F1, TP-3F2, TP-3G1, TP-3G3, TP-3H2, TP-4A7, TP-4A9, TP-4B7, TP-4E8, TP-4G8, or

[0137] The CDR1, 2, and/or 3 regions of the engineered antibodies described above can comprise the exact amino acid sequence(s) as those of TP-2A2, TP-2A5.1, TP-2A6, TP-2A8, TP-2A10, TP-2B1, TP-2B3, TP-2B4, TP-2B8, TP-2B9, TP-2C1, TP-2C7, TP-2D7, TP-2E3, TP-2E5, TP-2F9, TP-2G2, TP-2G4, TP-2G5, TP-2G6, TP-2G7, TP-2G9, TP-2H10, TP-3A2, TP-3A3, TP-3A4, TP-3B3, TP-3B4, TP-3C1, TP-3C2, TP-3C3, TP-3D3, TP-3E1, TP-3F1, TP-3F2, TP-3G1, TP-3G3, TP-3H2, TP-4A7, TP-4A9, TP-4B7, TP-4E8, TP-4G8, or TP-4H8 disclosed herein.

[0138] However, the ordinarily skilled artisan will appreciate that some deviation from the exact CDR sequences of TP-2A2, TP-2A5.1, TP-2A6, TP-2A8, TP-2A10, TP-2B1, TP-2B3, TP-2B4, TP-2B8, TP-2B9, TP-2C1, TP-2C7, TP-2D7, TP-2E3, TP-2E5, TP-2F9, TP-2G2, TP-2G4, TP-2G5, TP-2G6, TP-2G7, TP-2G9, TP-2H10, TP-3A2, TP-3A3, TP-3A4, TP-3B3, TP-3B4, TP-3C1, TP-3C2, TP-3C3, TP-3D3, TP-3E1, TP-3F1, TP-3F2, TP-3G1, TP-3G3, TP-3H2, TP-4A7, TP-4A9, TP-4B7, TP-4E8, TP-4G8, or TP-4H8 may be possible while still retaining the ability of the antibody to bind TFPI effectively. Accordingly, in another embodiment, the engineered antibody may be composed of one or more CDRs that are, for example, at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or 99.5% identical to one or more CDRs of TP-2A2, TP-2A5.1, TP-2A6, TP-2A8, TP-2A10, TP-2B1, TP-2B3, TP-2B4, TP-2B8, TP-2B9, TP-2C1, TP-2C7, TP-2D7, TP-2E3, TP-2E5, TP-2F9, TP-2G2, TP-2G4, TP-2G5, TP-2G6, TP-2G7, TP-2G9, TP-2H10, TP-3A2, TP-3A3, TP-3A4, TP-3B3, TP-3B4, TP-3C1, TP-3C2, TP-3C3, TP-3D3,

TP-3E1, TP-3F1, TP-3F2, TP-3G1, TP-3G3, TP-3H2, TP-4A7, TP-4A9, TP-4B7, TP-4E8, TP-4G8, or TP-4H8.

[0139] The antibody may be of any of the various classes of antibodies, such as without limitation an IgG1, an IgG2, an IgG3, an IgG4, an IgM, an IgA1, an IgA2, a secretory IgA, an IgD, and an IgE antibody.

[0140] In one embodiment, provided is an isolated fully human monoclonal antibody to human tissue factor pathway inhibitor.

[0141] In another embodiment, provided is an isolated fully human monoclonal antibody to Kunitz domain 2 of human tissue factor pathway inhibitor.

#### Nucleic Acids

[0142] Also provided are isolated nucleic acid molecules encoding any of the monoclonal antibodies described above.

#### Methods of Preparing Antibodies To TFPI

[0143] The monoclonal antibody may be produced recombinantly by expressing a nucleotide sequence encoding the variable regions of the monoclonal antibody according to the embodiments of the invention in a host cell. With the aid of an expression vector, a nucleic acid containing the nucleotide sequence may be transfected and expressed in a host cell suitable for the production. Accordingly, also provided is a method for producing a monoclonal antibody that binds with human TFPI comprising:

[0144] (a) transfecting a nucleic acid molecule encoding a monoclonal antibody of the invention into a host cell,
[0145] (b) culturing the host cell so to express the monoclonal antibody in the host cell, and optionally

[0146] (c) isolating and purifying the produced monoclonal antibody,

wherein the nucleic acid molecule comprises a nucleotide sequence encoding a monoclonal antibody of the present invention

[0147] In one example, to express the antibodies, or antibody fragments thereof, DNAs encoding partial or full-length light and heavy chains obtained by standard molecular biology techniques are inserted into expression vectors such that the genes are operatively linked to transcriptional and translational control sequences. In this context, the term "operatively linked" is intended to mean that an antibody gene is ligated into a vector such that transcriptional and translational control sequences within the vector serve their intended function of regulating the transcription and translation of the antibody gene. The expression vector and expression control sequences are chosen to be compatible with the expression host cell used. The antibody light chain gene and the antibody heavy chain gene can be inserted into separate vectors or, more typically, both genes are inserted into the same expression vector. The antibody genes are inserted into the expression vector by standard methods (e.g., ligation of complementary restriction sites on the antibody gene fragment and vector, or blunt end ligation if no restriction sites are present). The light and heavy chain variable regions of the antibodies described herein can be used to create full-length antibody genes of any antibody isotype by inserting them into expression vectors already encoding heavy chain constant and light chain constant regions of the desired isotype such that the  $\mathbf{V}_H$ segment is operatively linked to the C<sub>H</sub> segment(s) within the vector and the  $V_L$  segment is operatively linked to the  $C_L$ segment within the vector. Additionally or alternatively, the recombinant expression vector can encode a signal peptide that facilitates secretion of the antibody chain from a host cell. The antibody chain gene can be cloned into the vector such that the signal peptide is linked in-frame to the amino terminus of the antibody chain gene. The signal peptide can be an immunoglobulin signal peptide or a heterologous signal peptide (i.e., a signal peptide from a non-immunoglobulin protein).

[0148] In addition to the antibody chain encoding genes, the recombinant expression vectors of the invention carry regulatory sequences that control the expression of the antibody chain genes in a host cell. The term "regulatory sequence" is intended to include promoters, enhancers and other expression control elements (e.g., polyadenylation signals) that control the transcription or translation of the antibody chain genes. Such regulatory sequences are described, for example, in Goeddel; Gene Expression Technology. Methods in Enzymology 185, Academic Press, San Diego, Calif. (1990). It will be appreciated by those skilled in the art that the design of the expression vector, including the selection of regulatory sequences may depend on such factors as the choice of the host cell to be transformed, the level of expression of protein desired, etc. Examples of regulatory sequences for mammalian host cell expression include viral elements that direct high levels of protein expression in mammalian cells, such as promoters and/or enhancers derived from cytomegalovirus (CMV), Simian Virus 40 (SV40), adenovirus, (e.g., the adenovirus major late promoter (Ad-MLP)) and polyoma. Alternatively, nonviral regulatory sequences may be used, such as the ubiquitin promoter or β-globin promoter.

[0149] In addition to the antibody chain genes and regulatory sequences, the recombinant expression vectors may carry additional sequences, such as sequences that regulate replication of the vector in host cells (e.g., origins of replication) and selectable marker genes. The selectable marker gene facilitates selection of host cells into which the vector has been introduced (see, e.g., U.S. Pat. Nos. 4,399,216, 4,634,665 and 5,179,017, all by Axel et al.). For example, typically the selectable marker gene confers resistance to drugs, such as G418, hygromycin or methotrexate, on a host cell into which the vector has been introduced. Examples of selectable marker genes include the dihydrofolate reductase (DHFR) gene (for use in dhfr-host cells with methotrexate selection/amplification) and the neo gene (for G418 selection).

[0150] For expression of the light and heavy chains, the expression vector(s) encoding the heavy and light chains is transfected into a host cell by standard techniques. The various forms of the term "transfection" are intended to encompass a wide variety of techniques commonly used for the introduction of exogenous DNA into a prokaryotic or eukaryotic host cell, e.g., electroporation, calcium-phosphate precipitation, DEAE-dextran transfection and the like. Although it is theoretically possible to express the antibodies of the invention in either prokaryotic or eukaryotic host cells, expression of antibodies in eukaryotic cells, and most preferably mammalian host cells, is the most preferred because such eukaryotic cells, and in particular mammalian cells, are more likely than prokaryotic cells to assemble and secrete a properly folded and immunologically active antibody.

[0151] Examples of mammalian host cells for expressing the recombinant antibodies include Chinese Hamster Ovary (CHO cells) (including dhfr-CHO cells, described in Urlaub

and Chasin, (1980) Proc. Natl. Acad. Sci. USA 77:4216-4220, used with a DHFR selectable marker, e.g., as described in R. J. Kaufman and P. A. Sharp (1982) Mol. Biol. 159:601-621), NSO myeloma cells, COS cells, HKB11 cells and SP2 cells. When recombinant expression vectors encoding antibody genes are introduced into mammalian host cells, the antibodies are produced by culturing the host cells for a period of time sufficient to allow for expression of the antibody in the host cells or secretion of the antibody into the culture medium in which the host cells are grown. Antibodies can be recovered from the culture medium using standard protein purification methods, such as ultrafiltration, size exclusion chromatography, ion exchange chromatography and centrifugation.

#### Use of Partial Antibody Sequences To Express Intact Antibodies

[0152] Antibodies interact with target antigens predominantly through amino acid residues that are located in the six heavy and light chain CDRs. For this reason, the amino acid sequences within CDRs are more diverse between individual antibodies than sequences outside of CDRs. See, e.g., FIGS. 6 and 7, in which the CDR regions in the light and heavy variable chains, respectively, of the monoclonal antibody according to the present invention are identified. Because CDR sequences are responsible for most antibody-antigen interactions, it is possible to express recombinant antibodies that mimic the properties of specific naturally occurring antibodies by constructing expression vectors that include CDR sequences from the specific naturally occurring antibody grafted onto framework sequences from a different antibody with different properties (see, e.g., Riechmann, L. et al., 1998, Nature 332:323-327; Jones, P. et al., 1986, Nature 321:522-525; and Queen, C. et al., 1989, Proc. Natl. Acad. Sci. U.S.A. 86:10029-10033). Such framework sequences can be obtained from public DNA databases that include germline antibody gene sequences. These germline sequences will differ from mature antibody gene sequences because they will not include completely assembled variable genes, which are formed by V(D)J joining during B cell maturation. It is not necessary to obtain the entire DNA sequence of a particular antibody in order to recreate an intact recombinant antibody having binding properties similar to those of the original antibody (see WO 99/45962). Partial heavy and light chain sequence spanning the CDR regions is typically sufficient for this purpose. The partial sequence is used to determine which germline variable and joining gene segments contributed to the recombined antibody variable genes. The germline sequence is then used to fill in missing portions of the variable regions. Heavy and light chain leader sequences are cleaved during protein maturation and do not contribute to the properties of the final antibody. For this reason, it is necessary to use the corresponding germline leader sequence for expression constructs. To add missing sequences, cloned cDNA sequences can be combined with synthetic oligonucleotides by ligation or PCR amplification. Alternatively, the entire variable region can be synthesized as a set of short, overlapping, oligonucleotides and combined by PCR amplification to create an entirely synthetic variable region clone. This process has certain advantages such as elimination or inclusion or particular restriction sites, or optimization of particu-

[0153] The nucleotide sequences of heavy and light chain transcripts are used to design an overlapping set of synthetic

oligonucleotides to create synthetic V sequences with identical amino acid coding capacities as the natural sequences. The synthetic heavy and kappa chain sequences can differ from the natural sequences in three ways: strings of repeated nucleotide bases are interrupted to facilitate oligonucleotide synthesis and PCR amplification; optimal translation initiation sites are incorporated according to Kozak's rules (Kozak, 1991, J. Biol. Chem. 266:19867-19870); and HindIII sites are engineered upstream of the translation initiation sites.

[0154] For both the heavy and light chain variable regions, the optimized coding, and corresponding non-coding, strand sequences are broken down into 30-50 nucleotide sections at approximately the midpoint of the corresponding non-coding oligonucleotide. Thus, for each chain, the oligonucleotides can be assembled into overlapping double stranded sets that span segments of 150-400 nucleotides. The pools are then used as templates to produce PCR amplification products of 150-400 nucleotides. Typically, a single variable region oligonucleotide set will be broken down into two pools which are separately amplified to generate two overlapping PCR products. These overlapping products are then combined by PCR amplification to form the complete variable region. It may also be desirable to include an overlapping fragment of the heavy or light chain constant region in the PCR amplification to generate fragments that can easily be cloned into the expression vector constructs.

[0155] The reconstructed heavy and light chain variable regions are then combined with cloned promoter, translation initiation, constant region, 3' untranslated, polyadenylation, and transcription termination sequences to form expression vector constructs. The heavy and light chain expression constructs can be combined into a single vector, co-transfected, serially transfected, or separately transfected into host cells which are then fused to form a host cell expressing both chains.

**[0156]** Thus, in another aspect, the structural features of a human anti-TFPI antibody, e.g., TP2A8, TP2G6, TP2G7, TP4B7, etc., are used to create structurally related human anti-TFPI antibodies that retain the function of binding to TFPI. More specifically, one or more CDRs of the specifically identified heavy and light chain regions of the monoclonal antibodies of the invention can be combined recombinantly with known human framework regions and CDRs to create additional, recombinantly-engineered, human anti-TFPI antibodies of the invention.

[0157] Accordingly, in another embodiment, provided is a method for preparing an anti-TFPI antibody comprising: preparing an antibody comprising (1) human heavy chain framework regions and human heavy chain CDRs, wherein the human heavy chain CDR3 comprises an amino acid sequence selected from the amino acid sequences of SEQ ID NOs: 388-430 and/or (2) human light chain framework regions and human light chain CDRs, wherein the light chain CDR3 comprises an amino acid sequence selected from the amino acid sequences of SEQ ID NOs: 259-301; wherein the antibody retains the ability to bind to TFPI. In other embodiments, the method is practiced using other CDRs of the invention.

#### Pharmaceutical Compositions

[0158] Also provided are pharmaceutical compositions comprising therapeutically effective amounts of anti-TFPI monoclonal antibody and a pharmaceutically acceptable carrier. "Pharmaceutically acceptable carrier" is a substance that

may be added to the active ingredient to help formulate or stabilize the preparation and causes no significant adverse toxicological effects to the patient. Examples of such carriers are well known to those skilled in the art and include water, sugars such as maltose or sucrose, albumin, salts such as sodium chloride, etc. Other carriers are described for example in Remington's Pharmaceutical Sciences by E. W. Martin. Such compositions will contain a therapeutically effective amount of at least one anti-TFPI monoclonal antibody.

[0159] Pharmaceutically acceptable carriers include sterile aqueous solutions or dispersions and sterile powders for the extemporaneous preparation of sterile injectable solutions or dispersion. The use of such media and agents for pharmaceutically active substances is known in the art. The composition is preferably formulated for parenteral injection. The composition can be formulated as a solution, microemulsion, liposome, or other ordered structure suitable to high drug concentration. The carrier can be a solvent or dispersion medium containing, for example, water, ethanol, polyol (for example, glycerol, propylene glycol, and liquid polyethylene glycol, and the like), and suitable mixtures thereof. In some cases, it will include isotonic agents, for example, sugars, polyalcohols such as mannitol, sorbitol, or sodium chloride in the composition.

[0160] Sterile injectable solutions can be prepared by incorporating the active compound in the required amount in an appropriate solvent with one or a combination of ingredients enumerated above, as required, followed by sterilization microfiltration. Generally, dispersions are prepared by incorporating the active compound into a sterile vehicle that contains a basic dispersion medium and the required other ingredients from those enumerated above. In the case of sterile powders for the preparation of sterile injectable solutions, some methods of preparation are vacuum drying and freezedrying (lyophilization) that yield a powder of the active ingredient plus any additional desired ingredient from a previously sterile-filtered solution thereof.

#### Pharmaceutical Uses

[0161] The monoclonal antibody can be used for therapeutic purposes for treating genetic and acquired deficiencies or defects in coagulation. For example, the monoclonal antibodies in the embodiments described above may be used to block the interaction of TFPI with FXa, or to prevent TFPI-dependent inhibition of the TF/FVIIa activity. Additionally, the monoclonal antibody may also be used to restore the TF/FVIIa-driven generation of FXa to bypass the insufficiency of FVIII- or FIX-dependent amplification of FXa.

[0162] The monoclonal antibodies have therapeutic use in the treatment of disorders of hemostasis such as thrombocytopenia, platelet disorders and bleeding disorders (e.g., hemophilia A and hemophilia B). Such disorders may be treated by administering a therapeutically effective amount of the anti-TFPI monoclonal antibody to a patient in need thereof. The monoclonal antibodies also have therapeutic use in the treatment of uncontrolled bleeds in indications such as trauma and hemorrhagic stroke. Thus, also provided is a method for shortening the bleeding time comprising administering a therapeutically effective amount of an anti-TFPI monoclonal antibody of the invention to a patient in need thereof.

[0163] The antibodies can be used as monotherapy or in combination with other therapies to address a hemostatic disorder. For example, co-administration of one or more antibodies of the invention with a clotting factor such as factor

VIIa, factor VIII or factor IX is believed useful for treating hemophilia. In one embodiment, provided is a method for treating genetic and acquired deficiencies or defects in coagulation comprising administering (a) a first amount of a monoclonal antibody that binds to human tissue factor pathway inhibitor and (b) a second amount of factor VIII or factor IX, wherein said first and second amounts together are effective for treating said deficiencies or defects. In another embodiment, provided is a method for treating genetic and acquired deficiencies or defects in coagulation comprising administering (a) a first amount of a monoclonal antibody that binds to human tissue factor pathway inhibitor and (b) a second amount of factor VIII or factor IX, wherein said first and second amounts together are effective for treating said deficiencies or defects, and further wherein factor VII is not coadministered. The invention also includes a pharmaceutical composition comprising a therapeutically effective amount of the combination of a monoclonal antibody of the invention and factor VIII or factor IX, wherein the composition does not contain factor VII. "Factor VII" includes factor VII and factor VIIa. These combination therapies are likely to reduce the necessary infusion frequency of the clotting factor. By co-administration or combination therapy is meant administration of the two therapeutic drugs each formulated separately or formulated together in one composition, and, when formulated separately, administered either at approximately the same time or at different times, but over the same therapeutic period.

[0164] The pharmaceutical compositions may be parenterally administered to subjects suffering from hemophilia A or B at a dosage and frequency that may vary with the severity of the bleeding episode or, in the case of prophylactic therapy, may vary with the severity of the patient's clotting deficiency.

[0165] The compositions may be administered to patients in need as a bolus or by continuous infusion. For example, a bolus administration of an inventive antibody present as a Fab fragment may be in an amount of from 0.0025 to 100 mg/kg body weight, 0.025 to 0.25 mg/kg, 0.010 to 0.10 mg/kg or 0.10-0.50 mg/kg. For continuous infusion, an inventive antibody present as an Fab fragment may be administered at 0.001 to 100 mg/kg body weight/minute, 0.0125 to 1.25 mg/kg/min., 0.010 to 0.75 mg/kg/min., 0.010 to 1.0 mg/kg/ min. or 0.10-0.50 mg/kg/min. for a period of 1-24 hours, 1-12 hours, 2-12 hours, 6-12 hours, 2-8 hours, or 1-2 hours. For administration of an inventive antibody present as a fulllength antibody (with full constant regions), dosage amounts may be about 1-10 mg/kg body weight, 2-8 mg/kg, or 5-6 mg/kg. Such full-length antibodies would typically be administered by infusion extending for a period of thirty minutes to three hours. The frequency of the administration would depend upon the severity of the condition. Frequency could range from three times per week to once every two or three weeks.

[0166] Additionally, the compositions may be administered to patients via subcutaneous injection. For example, a dose of 10 to 100 mg anti-TFPI antibody can be administered to patients via subcutaneous injection weekly, biweekly or monthly.

[0167] As used herein, "therapeutically effective amount" means an amount of an anti-TFPI monoclonal antibody or of a combination of such antibody and factor VIII or factor IX that is needed to effectively increase the clotting time in vivo or otherwise cause a measurable benefit in vivo to a patient in need. The precise amount will depend upon numerous fac-

tors, including, but not limited to the components and physical characteristics of the therapeutic composition, intended patient population, individual patient considerations, and the like, and can readily be determined by one skilled in the art.

#### **EXAMPLES**

General Materials And Methods

Example 1

Panning And Screening of Human Antibody Library Against Human TFPI

Panning Human Antibody Library Against TFPI

[0168] Anti-TFPI antibodies were selected by panning phage displayed combinatorial human antibody library HuCal Gold (Rothe et al., J. Mol. Biol., 2008, 376: 1182-1200) against human TFPI (American Diagnostica). Briefly, 200 µl of TFPI (5 µg/ml) was coated on 96-well Maxisorp plates for overnight at 4° C. and the plates were then blocked with a PBS buffer containing 5% milk. After the plates were washed with PBS containing 0.01% Tween-20 (PBST), an aliquot of combinatorial human antibody library was added to the TFPI-coated wells and incubated for 2 hours. Unbound phage was washed away with PBST, and the antigen-bound phage was eluted with dithiothreitol, infected and amplified in *E. coli* strain TG1. The phage was rescued by helper phage for next round of panning. A total of three rounds of panning

were conducted and the clones from last two rounds were screened against human TFPI in an ELISA assay.

Screening Antibody Clones By Antigen-Binding In An ELISA

[0169] To select antibody clones that bind to human TFPI, Fab genes of the phage clones from the second and third round of panning were subcloned into a bacterial expression vector and expressed in *E. coli* strain TG1. The bacterial lysate was added to the wells of the human TFPI-coated Maxisorp plates. After washing, HRP-conjugated goat antihuman Fab was used as a detection antibody and the plates were developed by adding AmplexRed (Invitrogen) with hydrogen peroxide. A signal of at least five-fold higher than the background was considered as positive. The cross reactivity of the anti-human TFPI antibodies to mouse TFPI was determined by a similar mouse TFPI-binding ELISA. The plates were coated with mouse TFPI (R&D System), BSA and lysozyme. The later two antigens were used as negative controls. A representative set of data is shown in FIG. 1.

#### Sequences of Anti-TFPI Human Antibodies

[0170] After the panning and screening of the HuCal Gold human antibody library against TFPI, DNA sequencing was performed on the positive antibody clones, resulting in 44 unique antibody sequences (Table 2). Among these antibody sequences, 29 were lambda light chains and 15 were kappa light chains. Our analysis of variable region of heavy chains reveals 28 of VH3, 14 of VH6, 1 of VH1 and 1 of VH5.

TABLE 2

	Peptide sequence of variable region	n of 44 anti-TFPI antibodies
Clone	VL	VH
TP-2A2	DIELTQPPSVSVAPGQTARISCSGDNIRTYYVHWYQQKPGQ APVVVIYGDSKRPSGIPERFSGSNSGNTATLTISGTQAEDE ADYYCQSYDSEADSEVFGGGTKLTVLGQ (SEQ ID NO: 2)	QVQLVESGGGLVQPGGSLRLSCAASGFTFSNNAMNWVRQAP GKGLEWVSTISYDGSNTYYADSVKGRFTISRDNSKNTLYLQ MNSLRAEDTAVYYCARQAGGWTYSYTDVWGQGTLVTVSS (SEQ ID NO: 4)
TP-2A5.1	DIELTQPPSVSVAPGQTARISCSGDNIPEKYVHWYQQKPGQ APVLVIHGDNNRPSGIPERFSGSNSGNTATLTISGTQAEDE ADYYCQSFDAGSYFVFGGGTKLTVLGQ (SEQ ID NO: 6)	QVQLVESGGGLVQPGGSLRLSCAASGFTFSSYGSWVRQAPG KGLEWVSVISGSGSSTYYADSVKGRFTISRDNSKNTLYLQM NSLRAEDTAVYYCARVNISTHFDVWGQGTLVTVSS (SEQ ID NO: 8)
TP-2A6	DIELTQPPSVSVAPGQTARISCSGDKIGSKYVYWYQQKPGQ APVLVIYDSNRPSGIPERFSGSNSGNTATLTISGTQAEDEA DYYCASYDSIYSYWVFGGGTKLTVLGQ (SEQ ID NO: 10)	QVQLVESGGGLVQPGGSLRLSCAASGFTFSRYAMSWVRQAP GKGLEWVSSIISSSSETYYADSVKGRPTISRDNSKNTLYLQ MNSLRAEDTAVYYCARLMGYGHYYPFDYWGQGTLVTVSS (SEQ ID NO: 12)
TP-2A8	DIELTQPPSVSVAPGQTARISCSGDNLRNYYAHWYQQKPGQ APVVVIYYDNNRPSGIPERFSGSNSGNTATLTISGTQAEDE ADYYCQSWDDGVPVFGGGTKLTVLGQ (SEQ ID NO: 14)	QVQLVESGGGLVQPGGSLRLSCAASGFTFRSYGMSWVRQAP GKGLEWVSSIRGSSSSTYYADSVKGRFTISRDNSKNTLYLQ MNSLRAEDTAVYYCARKYRYWFDYWGQGTLVTVSS (SEQ ID NO: 16)
TP-2A10	DIELTQPPSVSVAPGQTARISCSGDKLGKKYVHWYQQKPGQ APVLVIYGDDKRPSGIPERFSGSNSGNTATLTISGTQAEDE ADYYCQAWGSISRFVFGGGTKLTVLGQ (SEQ ID NO: 18)	QVQLVESGGGLVQPGGSLRLSCAASGFTFTSYSMNWVRQAP GKGLEWVSAISYTGSNTHYADSVKGRFTISRDNSKNTLYLQ MNSLRAEDTAVYYCARAFLGYKESYFDIWGQGTLVTVSS (SEQ ID NO: 20)
TP-2B1	DIELTQPPSVSVAPGQTARISCSGDNLGNKYAHWYQQKPGQ APVLVIYYDNKRPSGIPERFSGSNSGNTATLTISGTQAEDE ADYYCQSWTPGSNTMVFGGGTRLTVLGQ (SEQ ID NO: 22)	QVQLVESGGGLVQPGGSLRLSCAASGFTFSSYSMSWVRQAS GKGLEWVSSIKGSGSNTYYADSVKGRPTISRDNSKNTLYLQ MNSLRAEDTAVYYCARNGGLIDVWGQGTLVTVSS (SEQ ID NO: 24)
TP-2B3	DIVLTQSPATLSLSPGERATLSCRASQNIGSNYLAWYQQKP GQAPRLLIYGASTRATGVPARFNGSGSGTDFTLTISSLEPE DFAVYYCQQLNSIPVTFGQGTKVEIKRT (SEQ ID NO: 26)	QVQLQQSGPGLVKPSQTLSLTCAISGDSVSSNSAAWGWIRQ SPGRGLEWLGMIYYRSKWYNSYAVSVKSRITINPDTSKNQF SLQLNSVTPEDTAVYYCARTMSKYGGPGMDVWGQGTLVTVS S (SEQ ID NO: 28)

TABLE 2-continued

	Peptide sequence of variable region	1 OI II GIICI IIII GIICIDOGICD
Clone	VL	VH
TP-2B4	DIELTQPPSVSVAPGQTARISCSGDALGTYYAYWYQQKPGQ APVLVIYGDMNRPSGIPERFSGSNSGNTATLTISGTQAEDE ADYYCQSYDAGVKPAVFGGGTKLTVLGQ (SEQ ID NO: 30)	QVQLVESGGGLVQPGGSLRLSCAASGFTFSNYSMTWVRQAP GKGLEWVSGISYNGSNTYYADSVKGRFTISRDNSKNTLYLQ MNSLRAEDTAVYYCARIYYMNLLAGWGQGTLVTVSS (SEQ ID NO: 32)
TP-2B8	DIELTQPPSVSVAPGQTARISCSGDNLRGYYASWYQQKPGQ APVLVIYEDNNRPSGIPERFSGSNSGNTATLTISGTQAEDE ADYYCQSWDSPYVHVFGGGTKLTVLGQ (SEQ ID NO: 34)	QVQLVQSGAEVKKPGASVKVSCKASGYTFTGNSMHWVRQAP GQGLEWMGTIFPYDGTTKYAQKFQGRVTMTRDTSISTAYME LSSLRSEDTAVYYCARGVHSYFDYWGQGTLVTVSS (SEQ ID NO: 36)
TP-2B9	DIQMTQSPSSLSASVGDRVTITCRASQSIRSYLAWYQQKPG KAPKLLIYKASNLQSGVPSRFSGSGSGTDFILTISSLQPED FAVYYCHQYSDSPVTFGQGTKVEIKRT (SEQ ID NO: 38)	QVQLQQSGPGLVKPSQTLSLTCAISGDSVSSNSAAWGWIRQ SPGRGLEWLGMIYHRSKWYNDYAVSVKSRITINPDTSKNQF SLQLNSVTPEDTAVYYCARYSSIGHMDYWGQGTLVTVSS (SEQ ID NO: 40)
TP-2C1	DIELTQPPSVSVAPGQTARISCSGDSIGSYYAHWYQQKPGQ APVLVIYYDSKRPSGIPERFSGSNSGNTATLTISGTQAEDE ADYYCQAYTGQSISRVFGGGTKLTVLGQ (SEQ ID NO: 42)	QVQLVESGGGLVQPGGSLRLSCAASGFTFSPYVMSWVRQAP GKGLEWVSSISSSSSNTYYADSVKGRFTISRDNSKNTLYLQ MNSLRAEDTAVYYCARGDSYMYDVWGQGTLVTVSS (SEQ ID NO: 44)
TP-2C7	DIQMTQSPSSLSASVGDRVTITCRASQDIRNNLAWYQQKPG KAPKLLIYAASSLQSGVPSRFSGSGSGTDFILTISSLQPED FAVYYCQQRNGFPLTFGQGTKVEIKRT (SEQ ID NO: 46)	QVQLQQSGPGLVKPSQTLSLTCAISGDSVSSNSAAWGWIRQ SPGRGLEWLGIIYYRSKWYNHYAVSVKSRITINPDTSKNQF SLQLNSVTPEDTAVYYCARSNWSGYFDYWGQGTLVTVSS (SEQ ID NO: 48)
TP-2D7	DIVMTQSPLSLPVTPGEPASISCRSSQSLLHSNGYTYLSWY LQKPGQSPQLLIYLGSNRASGVPDRFSGSGSGTDFTLKISR VEAEDVGVYYCQQYDNAPITEGQGTKVEIKRT (SEQ ID NO: 50)	QVQLQQSGPGLVKPSQTLSLTCAISGDSVSSNSAAWGWIRQ SPGRGLEWLGLIYYRSKWYNDYAVSVKSRITINPDTSKNQF SLQLNSVTPEDTAVYYCARFGDTNRNGTDVWGQGTLVTVSS (SEQ ID NO: 52)
TP-2E3	DIALTQPASVSGSPGQSITISCTGTSSDIGGYNYVSWYQQH PGKAPKLMIYGVNYRPSGVSNRFSGSKSGNTASLTISGLQA EDEADYYCSSADKFTMSIVFGGGTKLTVLGQ (SEQ ID NO: 54)	QVQLQQSGPGLVKPSQTLSLTCAISGDSVSSNSAAWGWIRQ SPGRGLEWLGMIYYRSKWYNDYAVSVKSRITINPDTSKNQF SLQLNSVTPEDTAVYYCARVNQYTSSDYWGQGTLVTVSS (SEQ ID NO: 56)
TP-2E5	DIQMTQSPSSLSASVGDRVTITCRASQPIYNSLSWYQQKPG KAPKLLIYGVSNLQSGVPSRFSGSGSGTDFILTISSLQPED FAVYYCLQVDNLPITFGQGTKVEIKRT (SEQ ID NO: 58)	QVQLQQSGPGLVKPSQTLSLTCAISGDSVSSNSAAWSWIRQ SPGRGLEWLGMIFYRSKWNNDYAVSVKSRITINPDTSKNQF SLQLNSVTPEDTAVYYCARVNANGYYAYVDLWGQGTLVTVS S
		(SEQ ID NO: 60)
TP-2F9	DIVLTQSPATLSLSPGERATLSCRASQSVSSQYLAWYQQKP GQAPRLLIYAASSRATGVPARFSGSGSGTDFTLTISSLEPE DFAVYYCQQDSNLPATFGQGTKVEIKRT (SEQ ID NO: 62)	QVQLVESGGGLVQPGGSLRLSCAASGFTFYKYAMHWVRQAP GKGLEWVSGIQYDGSYTYYADSVKGRFTISRDNSKNTLYLQ MNSLRAEDTAVYYCARYYCKCVDLWGQGTLVTVSS (SEQ ID NO: 64)
TP-2G2	DIELTQPPSVSVAPGQTARISCSGDNIRKFYVHWYQQKPGQ APVLVIYGTNKRPSGIPERFSGSNSGNTATLTISGTQAEDE ADYYCQSYDSKFNTVFGGGTKLTVLGQ (SEQ ID NO: 66)	QVQLVESGGGLVQPGGSLRLSCAASGFTFSSYAMNWVRQAP GKGLEWVSAILSDGSSTSMADSVKGRFTISRDNSKNTLYLQ MNSLRAEDTAVYYCARYPDWGWYTDVWGQGTLVTVSS (SEQ ID NO: 68)
TP-2G4	DIELTQPPSVSVAPGQTARISCSGDALRKHYVYWYQQKPGQ APVLVIYGDNNRPSGIPERFSGSNSGNTATLTISGTQAEDE ADYYCQSYDKPYPILVFGGGTKLTVLGQ (SEQ ID NO: 70)	QVQLVESGGGLVQPGGSLRLSCAASGFTFSSYAMTWVRQAP GKGLEWVSNISYSGSNTYYADSVKGRFTISRDNSKNTLYLQ MNSLRAEDTAVYYCARVGYYYGFDYWGQGTLVTVSS (SEQ ID NO: 72)
TP-2G5	DIVLTQSPATLSLSPGERATLSCRASQNVSSNYLAWYQQKP GQAPRLLIYDASNRATGVPARFSGSGSTDFTLTISSLEPE DFAVYYCQQFYDSPQTFGQGTKVEIKRT (SEQ ID NO: 74)	QVQLQQSGPGLVKPSQTLSLTCAISGDSVSSNSAAWSWIRQ SPGRGLEWLGFIYYRSKWYNDYAVSVKSRITINPDTSKNQF SLQLNSVTPEDTAVYYCARHNPDLGFDYWGQGTLVTVSS (SEQ ID NO: 76)
TP-2G6	DIVLTQSPATLSLSPGERATLSCRASQYVTSSYLAWYQQKP GQAPRLLIYGSSRATGVPARFSGSGSGTDFTLTISSLEPED FATYYCQQYSSSPITFGQGTKVEIKRT (SEQ ID NO: 78)	QVQLQQSGPGLVKPSQTLSLTCAISGDSVSSSSAAWSWIRQ SPGRGLEWLGIIYYRSKWYNDYAVSVKSRITINPDTSKNQF SLQLNSVTPEDTAVYYCARHSMVGFDVWGQGTLVTVSS (SEQ ID NO: 80)
TP-2G7	DIELTQPPSVSVAPGQTARISCSGDNLGTYYVHWYQQKPGQ APVLVIYGDNNRPSGIPERFSGSNSGNTATLTISGTQAEDE ADYYCQTYDSNNESIVFGGGTKLTVLGQ (SEQ ID NO: 82)	QVQLVESGGGLVQPGGSLRLSCAASGFTFNSYAMSWVRQAP GKGLEWVSNISSNSSNTYYADSVKGRFTISRDNSKNTLYLQ MNSLRAEDTAVYYCARKGGGEHGFFPSDIWGQGTLVTVSS (SEQ ID NO: 84)

TABLE 2-continued

	Peptide sequence of variable region	n of 44 anti-TFPI antibodies
Clone	VL	VH
TP-2G9	DIALTQPASVSGSPGQSITISCTGTSSDLGGFNTVSWYQQH PGKAPKLMIYSVSSRPSGVSNRFSGSKSGNTASLTISGLQA EDEADYYCQSYDLNNLVFGGGTKLTVLGQ (SEQ ID NO: 86)	QVQLVESGGGLVQPGGSLRLSCAASGFTFNSYAMTWVRQAP GKGLEWVSAIKSDGSNTYYADSVKGRFTISRDNSKNTLYLQ MNSLRAEDTAVYYCARNDSGWFDVWGQGTLVTVSS (SEQ ID NO: 88)
TP-2H10	DIVLTQSPATLSLSPGERATLSCRASQSVSSFYLAWYQQKP GQAPRLLIYGSSSRATGVPARFSGSGSGTDFTLTISSLEPE DFATYYCQQYDSTPSTFGQGTKVEIKRT (SEQ ID NO: 90)	QVQLQQSGPGLVKPSQTLSLTCAISGDSVSSNGAAWGWIRQ SPGRGLEWLGFIYRRSKWYNSYAVSVKSRITINPDTSKNQF SLQLNSVTPEDTAVYYCARQDGMGGMDSWGQGTLVTVSS (SEQ ID NO: 92)
TP-3A2	DIELTQPPSVSVAPGQTARISCSGDNIGSRYAYWYQQKPGQ APVVVIYDDSDRPSGIPERFSGSNSGNTATLTISGTQAEDE ADYYCAAYTFYARTVFGGGTKLTVLGQ (SEQ ID NO: 94)	QVQLVESGGGLVQPGGSLRLSCAASGFTFSNYYLSWVRQAP GKGLEWVSGISYNGSSTNYADSVKGRFTISRDNSKNTLYLQ MNSLRAEDTAVYYCARMWRYSLGADSWGQGTLVTVSS (SEQ ID NO: 96)
TP-3A3	DIELTQPPSVSVAPGQTARISCSGDNIGSKYVHWYQQKPGQ APVVVIYEDSDRPSGIPERFSGSNSGNTATLTISGTQAEDE ADYYCQSWDKSEGYVFGGGTKLTVLGQ (SEQ ID NO: 98)	QVQLVESGGGLVQPGGSLRLSCAASGFTFNNNAISWVRQAP GKGLEWVSAINSSSSSTSMADSVKGRFTISRDNSKNTLYLQ MNSLRAEDTAVYYCARGHHRGHSWASFIDYWGQGTLVTVSS (SEQ ID NO: 100)
TP-3A4	DIELTQPPSVSVAPGQTARISCSGDNLRDKYASWYQQKPGQ APVLVIYSKSERPSGIPERFSGSNSGNTATLTISGTQAEDE ADYYCSSYTLNPNLNYVFGGGTKLTVLGQ (SEQ ID NO: 102)	QVQLVESGGGLVQPGGSLRLSCAASGFTFSSYWMHWVRQAP GKGLEWVSSISYDSSNTYYADSVKGRFTISRDNSKNTLYLQ MNSLRAEDTAVYYCARYGGMDYWGQGTLVTVSS (SEQ ID NO: 104)
TP-3B3	DIELTQPASVSVAPGQTARISCSGDNLRSKYAHWYQQKPGQ APVLVIYGDNNRPSGIPERFSGSNSGNTATLTISGTQAEDE ADYYCSAYAMGSSPVFGGGTKLTVLGQ (SEQ ID NO: 106)	QVQLVESGGGLVQPGGSLRLSCAASGFTFSSYGMHWVRQAP GKGLEWVSNISYMGSNTNYADSVKGRFTISRDNSKNTLYLQ MNSLRAEDTAVYYCARGLFPGYFDYWGQGTLVTVSS (SEQ ID NO: 108)
TP-3B4	DIQMTQSPSSLSASVGDRVTITCRASQNISNYLNWYQQKPG KAPKLLIYGESSLQSGVPSRFSGSGSGTDFILTISSLQPED FAVYYCQQYGNNPTTFGQGTKVEIKRT (SEQ ID NO: 110)	QVQLQQSGPGLVKPSQTLSLTCAISGDSVSSNGAAWGWIRQ SPGRGLEWLGHIYYRSKWYNSYAVSVKSRITINPDTSKNQF SLQLNSVTPEDTAVYYCARWGGIHDGDIYFDYWGQGTLVTV SS (SEQ ID NO: 112)
TP-3C1	DIALTQPASVSGSPGQSITISCTGTSSDLGGFNTVSWYQQH PGKAPKLMIYSVSSRPSGVSNRFSGSKSGNTASLTISGLQA EDEADYYCQSYDLNNLVFGGGTKLTVLGQ (SEQ ID NO: 86)	QVQLVESGGGLVQPGGSLRLSCAASGFTFSSYSMHWVRQAP GKGLEWVSGISYSSSFTYYADSVKGRFTISRDNSKNTLYLQ MNSLRAEDTAVYYCARALGGGVDYWGQGTLVTVSS (SEQ ID NO: 136)
TP-3C2	DIQMTQSPSSLSASVGDRVTITCRASQSITNYLNWYQQKPG KAPKLLIYDVSNLQSGVPSRFSGSGSGTDFILTISSLQPED FAVYYCQQYSGYPLTFGQGTKVEIKRT (SEQ ID NO: 114)	QVQLQQSGPGLVKPSQTLSLTCAISGDSVSSSSAAWSWIRQ SPGRGLEWLGMIYYRSKWYNHYAVSVKSRITINPDTSKNQF SLQLNSVTPEDTAVYYCARGGSGVMDVWGQGTLVTVSS (SEQ ID NO: 116)
TP-3C3	DIQMTQSPSSLSASVGDRVTITCRASQSINPYLNWYQQKPG KAPKLLIYAASNLQSGVPSRFSGSGSGTDFILTISSLQPED FAVYYCQQLDNRSITFGQGTKVEIKRT (SEQ ID NO: 118)	QVQLQQSGPGLVKPSQTLSLTCAISGDSVSSNSAAWGWIRQ SPGRGLEWLGVIYYRSKWYNDYAVSVKSRITINPDTSKNQF SLQLNSVTPEDTAVYYCARARAKKSGGFDYWGQGTLVTVSS (SEQ ID NO: 120)
TP-3D3	DIELTQPPSVSVAPGQTARISCSGDSLGSKFAHWYQQKPGQ APVLVIYDDSNRPSGIPERFSGSNSGNTATLTISGTQAEDE ADYYCSTYTSRSHSYVFGGGTKLTVLGQ (SEQ ID NO: 122)	QVQLVESGGGLVQPGGSLRLSCAASGFTFSSYASWVRQAPG KGLEWVSGISGDGSNTHYADSVKGRFTISRDNSKNTLYLQM NSLRAEDTAVYYCARYDNFYFDVWGQGTLVTVSS (SEQ ID NO: 124)
TP-3E1	DIELTQPPSVSVAPGQTARISCSGDNIGSYYAYWYQQKPGQ APVLVIYDDSNRPSGIPERFSGSNSGNTATLTISGTQAEDE ADYYCQSYDSTGLLVFGGGTKLTVLGQ (SEQ ID NO: 126)	QVQLVESGGGLVQPGGSLRLSCAASGFTFSNYAMTWVRQAP GKGLEWVSVISSVGSNTYYADSVKGRFTISRDNSKNTLYLQ MNSLRAEDTAVYYCARPTKAGRTWWWGPYMDVWGQGTLVTV SS (SEQ ID NO: 128)
TP-3F1	DIELTQPPSVSVAPGQTARISCSGDNIGSYFASWYQQKPGQ APVLVIYDDSNRPSGIPERFSGSNSGNTATLTISGTQAEDE ADYYCEGSNVFGGGTKLTVLGQ (SEQ ID NO: 130)	QVQLVQSGAEVKKPGESLKISCKGSGYSFTDYWIGWVRQMP GKGLEWMGIIQPSDSDTNYSPSFQGQVTISADKSISTAYLQ WSSLKASDTAMYYCARFNWWGKYDSGEDVWGQGTLVIVSS (SEQ ID NO: 132)
TP-3F2	DIELTQPPSVSVAPGQTARISCSGDNLPSKSVYWYQQKPGQ APVLVIYGDNNRPSGIPERFSGSNSGNTATLTISGTQAEDE ADYYCQSWTSRPMVVFGGGTKLTVLGQ (SEQ ID NO: 134)	QVQLVESGGGLVQPGGSLRLSCAASGFTFSSYSMHWVRQAP GKGLEWVSGISYSSSFTYYADSVKGRFTISRDNSKNTLYLQ MNSLRAEDTAVYYCARALGGGVDYWGQGTLVTVSS (SEQ ID NO: 136)

TABLE 2-continued

	Peptide sequence of variable region	n of 44 anti-TFPI antibodies
Clone	VL	VH
TP-3G1	DIQMTQSPSSLSASVGDRVTITCRASQGISSYLHWYQQKPG KAPKLLIYGASTLQSGVPSRFSGSGSGTDFILTISSLQPED FATYYCQQQNGYPFTFGQGTKVEIKRT (SEQ ID NO: 138)	QVQLQQSGPGLVKPSQTLSLTCAISGDSVSSNSGGWGWIRQ SPGRGLEWLGLIYYRSKWYNAYAVSVKSRITINPDTSKNQF SLQLNSVTPEDTAVYYCARYLGSNFYVYSDVWGQGTLVTVS S (SEQ ID NO: 140)
TP-3G3	DIQMTQSPSSLSASVGDRVTITCRASQNIHSHLNWYQQKPG KAPKLLIYDASSLQSGVPSRFSGSGSGTDFILTISSLQPED FAVYYCQQYYDYPLTFGQGTKVEIKRT (SEQ ID NO: 142)	QVQLVESGGGLVQPGGSLRLSCAASGFTFSSYSMSWVRQAP GKGLEWVSSISSSSSNTYYGDSVKGRFTISRDNSKNTLYLQ MNSLRAEDTAVYYCARMHYKGMDIWGQGTLVTVSS (SEQ ID NO: 144)
TP-3H2	DIELTQPPSVSVAPGQTARISCSGDKLGKYYAYWYQQKPGQ APVLVIYGDSKRPSGIPERFSGSNSGNTATLTISGTQAEDE ADYYCSSAAFGSTVFGGGTKLTVLGQ (SEQ ID NO: 146)	QVQLVESGGGLVQPGGSLRLSCAASGFTFNSYYMSWVRQAP GKGLEWVSNISSSGSNTNYADSVKGRFTIGRDNSKNTLYLQ MNSLRAEDTAVYYCARVHYGFDFWGQGTLVTVSS (SEQ ID NO: 148)
TP-4A7	DIELTQPPSVSVAPGQTARISCSGDALGSKFAHWYQQKPGQ APVLVIYDDSERPSGIPERFSGSNSGNTATLTISGTQAEDE ADYYCQAYDSGLLYVFGGGTKLTVLGQ (SEQ ID NO: 150)	QVQLVESGGGLVQPGGSLRLSCAASGFTFRNYAMNWVRQAP GKGLEWVSVISGSSSYTYYADSVKGRFTISRDNSKNTLYLQ MNSLRAEDTAVYYCARADLPYMVFDYWGQGTLVTVSS (SEQ ID NO: 152)
TP-4A9	DIELTQPPSVSVAPGQTARISCSGDALGKYYASWYQQKPGQ APVLVIYGDNKRPSGIPERFSGSNSGNTATLTISGTQAEDE ADYYCQSYTTRSLVFGGGTKLTVLGQ (SEQ ID NO: 154)	QVQLVESGGGLVQPGGSLRLSCAASGFTFSSYGMSWVRQAP GKGLEWVSLISGVSSSTYYADSVKGRFTISRDNSKNTLYLQ MNSLRAEDTAVYYCARSYLGYFDVWGQGTLVTVSS (SEQ ID NO: 156)
TP-4B7	DIVMTQSPLSLPVTPGEPASISCRSSQSLVFSDGNTYLNWY LQKPQQSPQLLIYKGSNRASGVPDRFSGSGSGTDFTLKISR VEAEDVGVYYCQQYDSYPLITGQGTKVEIKRT (SEQ ID NO: 158)	QVQLQQSGPGLVKPSQTLSLTCAISGDSVSSNSAAWSWIRQ SPGRGLEWLGIIYKRSKWYMDYAVSVKSRITINPDTSKNQF SLQLNSVTPEDTAVYYCARWHSDKHWGFDYWGQGTLVTVSS (SEQ ID NO: 160)
TP-4E8	DIELTQPPSVSVAPGQTARISCSGDALGSKYVSWYQQKPGQ APVLVIYGDNKRPSGIPERFSGSNSGNTATLTISGTQAEDE ADYYCQSYTYSLNQVPGGGTKLTVLGQ (SEQ ID NO: 162)	QVQLVESGGGLVQPGGSLRLSCAASGFTFNDYAMSWVRQAP GKGLEWVSLIESVSSSTYYADSVKGRFTISRDNSKEELYLQ MNSLRAEDTAVYYCARTIGVLWDDVWGQGTLVTVSS (SEQ ID NO: 164)
TP-4G8	DIELTQPPSVSVAPGQTARISCSGDKLGSKSVHWYQQKPGQ APVLVIYRDTDRPSGIPERFSGSNSGNTATLTISGTQAEDE ADYYCQTYDYILNVFGGGTKLTVLGQ (SEQ ID NO: 166)	QVQLVESGGGLVQPGGSLRLSCAASGFTFSTYAMHWVRQAP GKGLEWVSTISGYGSFTYYADSVKGRFTISRDNSKEELYLQ MNSLRAEDTAVYYCARNGRKYGQMDNWGQGTLVTVSS (SEQ ID NO: 168)
TP-4H8	DIELTQPPSVSVAPGQTARISCSGDSIGKKYVHWYQQKPGQ APVLVIYGDNNRPSGIPERFSGSNSGNTATLTISGTQAEDE ADYYCSTADSVITYKNVFGGGTKLTVLGQ (SEQ ID NO: 170)	QVQLVESGGGLVQPGGSLRLSCAASGFTFSDHAMHWVRQAP GKGLEWVSVIEYSGSKTNYADSVKGRFTISRDNSKEELYLQ MNSLRAEDTAVYYCARGDYYPYLVFAIWGQGTLVTVSS (SEQ ID NO: 172)

### Cross-Reactivity To Mouse TFPI

[0171] The above 44 human TFPI-binding clones were also tested for binding to mouse TFPI in ELISA. Nineteen antibodies were found cross-reactive to mouse TFPI. To facilitate the study using mouse hemophilia model, we further characterized these 19 antibodies as well as five antibodies that were specific to human TFPI. A representative set of data is shown in FIG. 1. None of these antibodies bound to BSA or lysozyme in ELISA.

### Example 2

#### Expression And Purification of Anti-TFPI Antibodies

[0172] Anti-TFPI antibodies (as Fab fragments) were expressed and purified from the bacterial strain TG1. Briefly, a single colony of bacterial strain TG1 containing the antibody expression plasmid was picked and grown overnight in 8 ml of 2xYT medium in the presence of 34 µg/mlchloramphenicol and 1% glucose. A volume of 7 ml culture was

transferred to 250 ml fresh 2xYT medium containing 34  $\mu$ g/ml chloramphenicol and 0.1% glucose. After 3 hours of incubation, 0.5 mM IPTG was added to induce Fab expression. The culture was continued overnight at 25° C. The culture was centrifuged to pellet the bacterial cells. The pellet was then resuspended in a Bug Buster lysis buffer (Novagen). After centrifugation, the supernatant of bacterial lysis was filtered. The Fab fragments were affinity-purified through a Ni-NTA column (Qiagen) according to the manufacturer's instruction.

#### Example 3

#### Determination of $EC_{50}$ And Binding Affinity of Anti-TFPI Antibodies

[0173] Purified Fab antibodies were used to determine  $EC_{50}$  of anti-TFPI antibodies to human or mouse TFPI.  $EC_{50}$  was assessed in an ELISA, similarly as described above. The results were analyzed using SoftMax. The binding affinity of

anti-TFPI antibodies was determined in a Biacore assay. Briefly, the antigen, either human or mouse TFPI, was immobilized on the CM5-chips using the amine coupling kit (GE HealthCare) according to the instructions of the manufacturer. The amount of immobilized TFPI was adjusted to the mass of the antigen to give approximate 300 RU. The antibody Fabs were analyzed in mobile phase and at least five different concentrations (0.1, 0.4, 1.6, 6.4 and 25 nM) of the purified antibodies were used in the Biacore assay. The kinetics and binding affinity were calculated using Biacore T100 Evaluation software.

**[0174]** As shown in Table 3, the 24 anti-TFPI Fabs showed various  $EC_{50}$  to human TFPI (0.09 to 792 nM) and mouse TFPI (0.06 to 1035 nM), and the affinity determined by Biacore was accordingly various to human TFPI (1.25 to 1140 nM). In the Biacore study of the Fabs to mouse TFPI, the variation of affinity was smaller (3.08 to 51.8 nM).

TABLE 3

The binding activity of 24 antibodies against human or mouse TFPI as determined by ELISA and Biacore (hTFPI: human TFPI; mTFPI: mouse TFPI; Neg: signal was less than two fold of background; ND, not done).

	Binding E	C <sub>50</sub> (nM)	Affinit	y (nM)
Antibody clones	hTFPI	mTFPI	hTFPI	mTFPI
TP-2A2	0.62	1035.88	6.57	29.8
TP-2A5	28.64	14.54	35.4	19.6
TP-2A8	0.09	0.06	1.25	3.08
TP-2B11	11.52	0.52	21.5	16.3
TP-2B3	0.84	20.18	7.40	27.0
TP-2C1	0.40	Neg	2.64	Neg
TP-2C7	0.60	0.60	2.01	9.33
TP-2E5	791.60	202.28	115	25.2
TP-2G5	342.52	871.34	42.1	16.1
TP-2G6	0.48	5.18	5.06	46.1
TP-2G7	23.48	Neg	26.9	Neg
TP-2G9	10.80	194.42	48.5	35.7
TP-2H10	2.18	32.40	10.2	11.5
TP-3A4	42.84	326.58	21.6	23.7
TP-3B4	35.76	34.62	14.1	20.4
TP-3C1	32.80	108.40	21.6	33.6
TP-3C2	59.00	956.68	17.1	28.5
TP-3G1	74.40	8.68	1140	49.1
TP-3G3	33.60	47.06	16.0	25.7
TP-4A9	0.17	117.68	7.60	Neg
TP-4B7	0.74	2.64	15.8	51.8
TP-4E8	36.94	Neg	35.9	ND
TP-4G8	846.92	Neg	25.2	ND
TP-4H8	72.50	Neg	32.2	ND

# Example 4

# Conversion of Anti-TFPI Fab To IgG

[0175] All of the identified anti-TFPI antibodies are fully human Fabs that can be feasibly converted to human IgG as therapeutic agent. In this example, however, the selected Fabs were converted to a chimeric antibody containing a mouse IgG constant region, so they are more suitable for testing in mouse model. The variable region of the selected antibodies was grafted into a mammalian expression vector containing mouse constant regions. The fully assembled IgG molecule was then transfected and expressed in HKB11 cells (Mei et al., Mol. Biotechnol., 2006, 34: 165-178). The culture supernatant was collected and concentrated. The anti-TFPI IgG

molecules were affinity purified through a Hitrap Protein G column (GE Healthcare) following the manufacturer's instruction.

#### Example 5

#### Selection of Anti-TFPI Neutralizing Antibodies

[0176] Anti-TFPI neutralizing antibodies were selected based on their inhibition of the TFPI activity under three experimental conditions. The activity of TFPI was measured using ACTICHROME® TFPI activity assay (American Diagnostica Inc., Stamford, Conn.), a three stage chromogenic assay to measures the ability of TFPI to inhibit the catalytic activity of the TF/FVIIa complex to activate factor X to factor Xa. The neutralizing activity of the anti-TFPI antibody is proportional to the amount of the restored FXa generation. In the first setting, purified anti-TFPI antibodies were incubated with human or mouse recombinant TFPI (R&D System) at the indicated concentrations. After incubation, the samples were mixed with TF/FVIIa and FX, and the residual activity of the TF/FVIIa complex was then measured using SPECTROZYME® FXa, a highly specific fXa chromogenic substrate. This substrate was cleaved only by FXa generated in the assay, releasing a p-nitroaniline (pNA) chromophore, which was measured at 405 nm. The TFPI activity present in the sample was interpolated from a standard curve constructed using known TFPI activity levels. The assay was performed in an end-point mode. In two other settings, anti-TFPI antibodies were spiked into normal human plasma or hemophilic A plasma, and the restored FXa generation was measured.

#### Example 6

#### Anti-TFPI Antibodies Shorten Clotting Time In A Diluted Prothrombin Time (dPT) Assay

[0177] The dPT assay was carried out essentially as described in Welsch et al., Thrombosis Res., 1991, 64 (2): 213-222. Briefly, human normal plasma (FACT, George King Biomedical), human TFPI depleted plasma (American Diagnostica) or hemophilic A plasma (George King Biomedical) were prepared by mixing plasma with 0.1 volumes of control buffer or anti-TFPI antibodies. After incubation for 30 min at 25° C., plasma samples (100  $\mu$ l) were combined with 200  $\mu$ l of appropriately diluted (1:500 dilution) Simplastin (Biometieux) as a source of thromboplastin and the clotting time was determined using a fibrometer STA4 (Stago). Thromboplastin was diluted with PBS or 0.05 M Tris based buffer (pH 7.5) containing 0.1 M sodium chloride, 0.1% bovine serum albumin and 20  $\mu$ M calcium chloride.

#### Example 7

Anti-TFPI Antibodies, Alone Or In Combination With Recombinant Factor VIII Or Factor IX, Shorten Blood Clotting Time In A ROTEM Assay

[0178] The ROTEM system (Pentapharm GmbH) included a four-channel instrument, a computer, plasma standards, activators and disposable cups and pins. Thrombelastographic parameters of ROTEM hemostasis systems included: Clotting Time (CT), which reflects the reaction time (the time required to obtain 2 mm amplitude following the initiation of data collection) to initiate blood clotting; Clot Formation Time (CFT) and the alpha angle to reflect clotting propaga-

tion, and the maximum amplitude and the maximum elastic modulus to reflect clot firmness. In the ROTEM assay,  $300\,\mu l$  of fresh citrated whole blood or plasma was assessed. All constituents were reconstituted and mixed according to the manufacturer's instructions, with data collection for the time period required for each system. Briefly, samples were mixed by withdrawing/dispensing  $300\,\mu l$  of blood or plasma with an automated pipette into ROTEM cups with  $20\,\mu l$  of CaCl $_2$  (200 mmol) added, followed immediately by mixing of the sample and initiation of data collection. Data were collected for 2 hr using a computer-controlled (software version 2.96) ROTEM system.

[0179] An exemplary result of ROTEM assay in detecting the effect of anti-TFPI antibodies in shortening blood clotting time is shown in FIGS. 3 and 5. FIG. 3 shows that TP-2A8-Fab shortened clotting time in human hemophiliac A plasma or mouse hemophiliac A whole blood, alone or in combination with recombinant FVIII, when ROTEM system was initiated with NATEM. FIG. 5 shows that anti-TFPI antibodies in IgG format (TP-2A8, TP-3G1, and TP-3C2) shortened clotting times as compared to a negative control mouse IgG antibody. Based on these results and the understanding in the field, the skilled person would expect that these anti-TFPI antibodies also shorten clotting time in combination with recombinant FIX as compared to these antibodies alone.

#### Example 8

#### In Vitro Functional Activity of Anti-TFPI Antibodies

[0180] To investigate the TFPI antibodies in blocking the function of TFPI, both chromogenic assay ACTICHROME and diluted prothrombin time (dPT) were used to test the functional activity of the antibodies obtained from the panning and screening. In both assays, a monoclonal rat anti-TFPI antibody (R&D System) was used as positive control and human polyclonal Fab was used as negative control. In the chromogenic assay, eight of the antibodies inhibited more than 50% of TFPI activity compared with the rat monoclonal antibody (Table 4). In dPT assay, all of these eight anti-TFPI Fabs showed a highly inhibitory effect, shortening the clotting time below 80 seconds, and four of the eight Fabs shortened dPT below 70 seconds. Dose-dependence of four of representative clones in shortening the dPT is shown in FIG. 2. However, other human anti-TFPI Fabs with low or no TFPI inhibitory activity also shortened clotting time in dPT. For example, TP-3B4 and TP-2C7, although showing less than 25% inhibitory activity, could shorten the dPT to less than 70 seconds. A simple linear regression analysis of inhibitory activity and dPT suggests significant correlation (p=0.0095) but large variance (R square=0.258).

TABLE 4

The in vitro functional activity of the anti-TFPI

	in human TFPI assay and o	
clone	% inhibition of hTFPI activity	dPT in hemoA plasma (sec)
anti-TFPI	100%	63.5
TP-2B3	100%	74.0
TP-4B7	100%	53.9
TP-3G1	93%	75.1
TP-3C2	92%	68.9
TP-2G6	86%	62.8
TP-2A8	100%	57.9

TABLE 4-continued

The in vitro functional activity of the anti-TFPI antibodies as determined by their inhibition activity in human TFPI assay and dPT assay.

clone	% inhibition of hTFPI activity	dPT in hemoA plasma (sec)
TP-2H10	63%	79.5
TP-2G7	55%	72.2
TP-4G8	39%	73.9
TP-2G5	36%	73.2
TP-2A5	30%	70.8
TP-4E8	29%	71.9
TP-4H8	28%	76.5
TP-3B4	25%	69.1
TP-2A2	23%	70.9
TP-2C1	21%	70.9
TP-3G3	15%	70.7
TP-2E5	0%	79.0
TP-3A4	0%	72.3
TP-3C1	0%	72.3
TP-2B11	0%	82.6
TP-2C7	0%	62.5
TP-2G9	0%	82.7
Untreated	0%	92.9

[0181] One of the anti-TFPI Fab, Fab-2A8, was also tested in ROTEM assay in which either human hemophilia A plasma with a low level of factor VIII or mouse hemophilia A whole blood was used. As shown in FIG. 3, comparing a polyclonal rabbit anti-TFPI antibody, Fab-2A8 showed similar activity in human hemophilia A plasma, decreasing clotting time (CT) from 2200 seconds to approximate 1700 seconds. When mouse hemophilia A whole blood was used, the control antibody, rabbit anti-TFPI shortened CT from 2700 seconds to 1000 seconds, whereas Fab-2A8 shorten CT from 2650 seconds to 1700 seconds. These results indicate that Fab-2A8 can significantly shorten clotting time in both human plasma and mouse blood (p=0.03).

## Example 9

# Function of Anti-TFPI Antibodies Following Conversion To Chimeric IgG

**[0182]** In-vitro assays of factor Xa generation and diluted prothrombin time indicate that at least six of the 24 anti-TFPI Fabs, TP-2A8, TP-2B3, TP-2G6, TP-3C2, TP-3G1 and TP-4B7, could block TFPI function. To facilitate in vivo study using hemophilia A mice, we converted these six anti-TFPI human Fabs into chimeric IgG, using the murine IgG1 isotype. The IgG expression vector was transfected into HKB11 cells, and the expressed antibody was collected in the culture supernatant and purified on Protein G column. When a representative clone 2G6-Fab was converted to IgG, the 2G6-IgG showed two fold increase of  $EC_{50}$  binding to human TFPI (from 0.48 nM to 0.22 nM) and 10-fold increase to mouse TFPI (from 5.18 nM to 0.51 nM). The results of IgG-2G6 binding to human and mouse TFPI are shown in FIG. **4**.

#### Example 10

#### Effect On Survival Rate In Hemophilia A (HemA) Mouse Tail Vein Transection Model

[0183] A mouse tail vein transection model has been established for pharmacologic evaluation. This model simulates the wide range of bleeding phenotypes observed between

normal individuals and severe hemophiliacs. For these studies, male hemophilia A mice (8 weeks old and 20 to 26 grams) were used. Mice were dosed via tail vein infusion with anti-TFPI monoclonal antibody (40  $\mu$ g/mouse), alone or together with a clotting factor such as FVIII (0.1 IU/mouse) prior to the injury. At 24 hours post-dosing, the left vein of the tail at 2.7 mm from the tip (in diameter) was transected. Survival was observed over 24 hours post transection. Survival rate was demonstrated to be dose-dependent when given with recombinant FVIII (data not shown). Data shown in FIG. **8** were from two separate studies (n=15 and n=10, respectively). The results showed that TP-2A8-IgG significantly prolonged the survival of hemophilia A mice as compared to control mouse IgG; and, in combination with recombinant FVIII, displayed a better survival rate than either agent alone.

#### Example 11

Combination of Anti-TFPI Antibody With Recombinant Factor VIIa Further Shortened Clotting Time And Clot Formation Time

[0184] The combined effect of anti-TFPI antibody and recombinant FVIIa (Novo Nordisk) was assessed in a ROTEM system using EXTEM (1:1000 dilution) and mouse hemophilia A whole blood. The indicated amounts of anti-TFPI antibody, TP-2A8-IgG ("2A8"), and recombinant FVIIa ("FVIIa"), were added into 300 µl of citrated mouse hemophilia A whole blood, and blood clotting was initiated using EXTEM system. FIG. 9 shows that addition of TP-2A8-IgG or recombinant FVIIa into mouse hemophilia A whole blood shortened clotting time and clot formation time, respectively. Combination of TP-2A8-IgG and recombinant FVIIa ("2A8+FVIIa") further shortened clotting time and clot formation time, indicating that combination of anti-TFPI antibody with recombinant FVIIa is useful in the treatment of hemophilia patients with or without inhibitors.

#### Example 12

Anti-TFPI Antibodies Shortened Clotting Time In FVIII Inhibitor-Induced Human Hemophiliac Blood

[0185] Selected anti-TFPI antibodies, 2A8 and 4B7 were also tested in a ROTEM assay using neutralizing FVIII antibodies to induce hemophilia in whole blood drawn from non-hemophilic patienst. FIG. 10 shows that normal human blood has a clotting time of approximately 1000 seconds. In the presence of FVIII neutralizing antibodies (PAH, 100 microgram/mL), the clotting time was prolonged to approximately 5200 seconds. The prolonged clotting time was sig-

nificantly shortened by addition of an anti-TFPI antibody, 2A8 or 4B7, indicating that anti-TFPI antibody is useful in the treatment of hemophilia patients with inhibitors.

### Example 13

#### Inhibitory Anti-TFPI Antibodies Bind To Kunitz Domain 2 of Human TFPI

[0186] Western blots and ELISA were used to determine which domain(s) of TFPI of the inhibitory antibodies bind. Recombinant full length human TFPI or TFPI domains were used for these studies. ELISA was similar to Example 3. In the Western Blot, human TFPI or domains were run on 4-12% Bis-Tris SDS PAGE running buffer MES (Invitrogen, Carlsbad, Calif.) and then transferred to cellulose membrane. After incubation with inhibitory antibodies for 10 min, the membrane was washed three times using SNAPid system (Millipore, Billerica, Mass.). A HRP conjugated donkey antimouse antibody (Pierce, Rockford, Ill.) at 1 to 10,000 dilution was incubated with the membrane for 10 min. After a similar wash step, the membrane was developed using SuperSignal substrate (Pierce, Rockford, Ill.). Whereas the control anti-Kunitz domain 1 antibody binds to full length TFPI, truncated TFPI and domains, inhibitory anti-TFPI antibodies only bind to TFPI containing Kunitz domain 2. This indicates that binding to Kunitz domain 2 is necessary for antibody's inhibitory function.

TABLE 5

T		ins bour by West				ermine	i	
	Anti- K1	mlgG	TP- 2A8	TP- 2B3	TP- 2G6		TP- 3G1	TP- 4B7
Full length	+	_	+	+	+	+	+	+
K1 + K2 + K3	+	_	+	+	+	+	+	+
K1 + K2	+	_	+	+	+	+	+	+
K1	+	-	-	-	-	-	-	-

[0187] While the present invention has been described with reference to the specific embodiments and examples, it should be understood that various modifications and changes may be made and equivalents may be substituted without departing from the true spirit and scope of the invention. The specification and examples are, accordingly, to be regarded in an illustrative rather then a restrictive sense. Furthermore, all articles, books, patent applications and patents referred to herein are incorporated herein by reference in their entireties.

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agetgegegg ceteeggatt tacetttact tettatteta tgaattgggt gegeeaagee cctqqqaaqq qtctcqaqtq qqtqaqcqct atctcttata ctqqtaqcaa tacccattat 180 geggatageg tgaaaggeeg ttttaccatt teaegtgata attegaaaaa caecetgtat 240 ctgcaaatga acagcctgcg tgcggaagat acggccgtgt attattgcgc gcgtgctttt 300 cttggttata aggagtctta ttttgatatt tggggccaag gcaccctggt gacggttagc 360 tcagc 365 <210> SEQ ID NO 20 <211> LENGTH: 121 <212> TYPE: PRT <213 > ORGANISM: Homo sapiens <400> SEQUENCE: 20 Gln Val Gln Leu Val Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Thr Ser Tyr Ser Met Asn Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val Ser Ala Ile Ser Tyr Thr Gly Ser Asn Thr His Tyr Ala Asp Ser Val Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys 85 90 95 Ala Arg Ala Phe Leu Gly Tyr Lys Glu Ser Tyr Phe Asp Ile Trp Gly Gln Gly Thr Leu Val Thr Val Ser Ser <210> SEQ ID NO 21 <211> LENGTH: 330 <212> TYPE: DNA <213> ORGANISM: Homo sapiens <400> SEOUENCE: 21 gatategaac tgacceagee geetteagtg agegttgeac caggteagae egegegtate 60 tcgtgtagcg gcgataatct tggtaataag tatgctcatt ggtaccagca gaaacccggg 120 caggegeeag ttettgtgat ttattatgat aataagegte ceteaggeat eeeggaaege 180 tttagcggat ccaacagcgg caacaccgcg accctgacca ttagcggcac tcaggcggaa gacgaagcgg attattattg ccagtcttgg actcctggtt ctaatactat ggtgtttggc 300 ggcggcacga ggttaaccgt tcttggccag 330 <210> SEQ ID NO 22 <211> LENGTH: 110 <212> TYPE: PRT <213 > ORGANISM: Homo sapiens <400> SEQUENCE: 22 Asp Ile Glu Leu Thr Gln Pro Pro Ser Val Ser Val Ala Pro Gly Gln

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Thr Ala Arg Ile Ser Cys Ser Gly Asp Asn Leu Gly Asn Lys Tyr Ala
His Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Val Leu Val Ile Tyr
Tyr Asp Asn Lys Arg Pro Ser Gly Ile Pro Glu Arg Phe Ser Gly Ser
Asn Ser Gly Asn Thr Ala Thr Leu Thr Ile Ser Gly Thr Gln Ala Glu
Asp Glu Ala Asp Tyr Tyr Cys Gln Ser Trp Thr Pro Gly Ser Asn Thr
Met Val Phe Gly Gly Gly Thr Arg Leu Thr Val Leu Gly Gln
<210> SEQ ID NO 23
<211> LENGTH: 350
<212> TYPE: DNA
<213 > ORGANISM: Homo sapiens
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tctgggaagg gtctcgagtg ggtgagctct atcaagggtt ctggtagcaa tacctattat
gcggatagcg tgaaaggccg ttttaccatt tcacgtgata attcgaaaaa caccctgtat
ctgcaaatga acagcctgcg tgcggaagat acggccgtgt attattgcgc gcgtaatggt
ggtcttattg atgtttgggg ccaaggcacc ctggtgacgg ttagctcagc
<210> SEQ ID NO 24
<211> LENGTH: 116
<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 24
Gln Val Gln Leu Val Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Ser Tyr
Ser Met Ser Trp Val Arg Gln Ala Ser Gly Lys Gly Leu Glu Trp Val
Ser Ser Ile Lys Gly Ser Gly Ser Asn Thr Tyr Tyr Ala Asp Ser Val
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys
Ala Arg Asn Gly Gly Leu Ile Asp Val Trp Gly Gln Gly Thr Leu Val
Thr Val Ser Ser
       115
<210> SEQ ID NO 25
<211> LENGTH: 330
<212> TYPE: DNA
<213 > ORGANISM: Homo sapiens
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ctgagctgca gagcgagcca gaatattggt tctaattatc tggcttggta ccagcagaaa	120
ccaggtcaag caccgcgtct attaatttat ggtgcttcta ctcgtgcaac tggggtcccg	180
gcgcgtttta acggctctgg atccggcacg gattttaccc tgaccattag cagcctggaa	240
cctgaagact ttgcggttta ttattgccag cagcttaatt ctattcctgt tacctttggc	300
cagggtacga aagttgaaat taaacgtacg	330
<210> SEQ ID NO 26 <211> LENGTH: 110 <212> TYPE: PRT <213> ORGANISM: Homo sapiens	
<400> SEQUENCE: 26	
Asp Ile Val Leu Thr Gln Ser Pro Ala Thr Leu Ser Leu Ser Pro Gly 1 10 15	
Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Asn Ile Gly Ser Asn 20 25 30	
Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu 35 40 45	
Ile Tyr Gly Ala Ser Thr Arg Ala Thr Gly Val Pro Ala Arg Phe Asn 50 55 60	
Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Glu	
65 70 75 80	
Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Leu Asn Ser Ile Pro 85 90 95	
Val Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr 100 105 110	
<210> SEQ ID NO 27 <211> LENGTH: 374 <212> TYPE: DNA <213> ORGANISM: Homo sapiens	
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acctgtgcga tttccggaga tagcgtgagc tctaattctg ctgcttgggg ttggattcgc	120
cagteteetg ggegtggeet egagtggetg ggeatgatet attategtag caagtggtat	180
aactettatg eggtgagegt gaaaageegg attaceatea acceggatae ttegaaaaac	240
cagtttagec tgcaactgaa cagegtgaec eeggaagata eggeegtgta ttattgegeg	300
cgtactatgt ctaagtatgg tggtcctggt atggatgttt ggggccaagg caccctggtg	360
acggttaget cage	374
<210> SEQ ID NO 28 <211> LENGTH: 124 <212> TYPE: PRT <213> ORGANISM: Homo sapiens	
<400> SEQUENCE: 28	
Gln Val Gln Leu Gln Gln Ser Gly Pro Gly Leu Val Lys Pro Ser Gln 1 10 15	

<213 > ORGANISM: Homo sapiens

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Thr Leu Ser Leu Thr Cys Ala Ile Ser Gly Asp Ser Val Ser Ser Asn
Ser Ala Ala Trp Gly Trp Ile Arg Gln Ser Pro Gly Arg Gly Leu Glu
Trp Leu Gly Met Ile Tyr Tyr Arg Ser Lys Trp Tyr Asn Ser Tyr Ala
Val Ser Val Lys Ser Arg Ile Thr Ile Asn Pro Asp Thr Ser Lys Asn
Gln Phe Ser Leu Gln Leu Asn Ser Val Thr Pro Glu Asp Thr Ala Val
Tyr Tyr Cys Ala Arg Thr Met Ser Lys Tyr Gly Gly Pro Gly Met Asp
Val Trp Gly Gln Gly Thr Leu Val Thr Val Ser Ser
       115
<210> SEQ ID NO 29
<211> LENGTH: 330
<212> TYPE: DNA
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 29
gatategaac tgacceagec geetteagtg agegttgeac caggteagac egegegtate
tcgtgtagcg gcgatgctct tggtacttat tatgcttatt ggtaccagca gaaacccggg
caggegecag ttettgtgat ttatggtgat atgaategte ceteaggeat eeeggaacge
tttageggat ceaacagegg caacacegeg accetgacea ttageggeac teaggeggaa
                                                                     240
gacgaagegg attattattg ccagtettat gatgetggtg ttaageetge tgtgtttgge
                                                                     300
ggcggcacga agttaaccgt tcttggccag
                                                                     330
<210> SEQ ID NO 30
<211> LENGTH: 110
<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 30
Asp Ile Glu Leu Thr Gln Pro Pro Ser Val Ser Val Ala Pro Gly Gln
Thr Ala Arg Ile Ser Cys Ser Gly Asp Ala Leu Gly Thr Tyr Tyr Ala
Tyr Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Val Leu Val Ile Tyr
Gly Asp Met Asn Arg Pro Ser Gly Ile Pro Glu Arg Phe Ser Gly Ser
Asn Ser Gly Asn Thr Ala Thr Leu Thr Ile Ser Gly Thr Gln Ala Glu
Asp Glu Ala Asp Tyr Tyr Cys Gln Ser Tyr Asp Ala Gly Val Lys Pro
Ala Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu Gly Gln
                               105
<210> SEQ ID NO 31
<211> LENGTH: 355
<212> TYPE: DNA
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<400> SEOUENCE: 31
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gctgcgcggc ctccggattt accttttcta attattctat gacttgggtg cgccaagccc
                                                                      120
ctgggaaggg tctcgagtgg gtgagcggta tctcttataa tggtagcaat acctattatg
                                                                      180
cggatagcgt gaaaggccgt tttaccattt cacgtgataa ttcgaaaaac accctgtatc
                                                                      240
tgcaaatgaa cagcctgcgt gcggaagata cggccgtgta ttattgcgcg cgtatttatt
                                                                      300
atatgaatct tcttgctggt tggggccaag gcaccctggt gacggttagc tcagc
                                                                      355
<210> SEQ ID NO 32
<211> LENGTH: 118
<212> TYPE: PRT
<213> ORGANISM: Homo sapiens
<400> SEQUENCE: 32
Gln Val Gln Leu Val Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
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Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Asn Tyr
Ser Met Thr Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
Ser Gly Ile Ser Tyr Asn Gly Ser Asn Thr Tyr Tyr Ala Asp Ser Val
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys
Ala Arg Ile Tyr Tyr Met Asn Leu Leu Ala Gly Trp Gly Gln Gly Thr
                                105
            100
Leu Val Thr Val Ser Ser
       115
<210> SEQ ID NO 33
<211> LENGTH: 327
<212> TYPE: DNA
<213> ORGANISM: Homo sapiens
<400> SEQUENCE: 33
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tegtgtageg gegataatet tegtggttat tatgettett ggtaceagea gaaaceeggg
                                                                      120
caggegecag ttettgtgat ttatgaggat aataategte eetcaggeat eeeggaacge
tttageggat ccaacagegg caacacegeg accetgacea ttageggeac tcaggeggaa
                                                                      240
gacgaagegg attattattg ceagtettgg gatteteett atgtteatgt gtttggegge
                                                                      300
ggcacgaagt taaccgttct tggccag
                                                                      327
<210> SEQ ID NO 34
<211> LENGTH: 109
<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
Asp Ile Glu Leu Thr Gln Pro Pro Ser Val Ser Val Ala Pro Gly Gln
              5
                                 10
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Ser Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Val Leu Val Ile Tyr
                           40
Glu Asp Asn Asn Arg Pro Ser Gly Ile Pro Glu Arg Phe Ser Gly Ser
Asn Ser Gly Asn Thr Ala Thr Leu Thr Ile Ser Gly Thr Gln Ala Glu
Asp Glu Ala Asp Tyr Tyr Cys Gln Ser Trp Asp Ser Pro Tyr Val His
Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu Gly Gln
           100
<210> SEQ ID NO 35
<211> LENGTH: 353
<212> TYPE: DNA
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 35
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agctgcaaag cctccggata tacctttact ggtaattcta tgcattgggt ccgccaagcc
cctqqqcaqq qtctcqaqtq qatqqqcact atctttccqt atqatqqcac tacqaaqtac
qcqcaqaaqt ttcaqqqccq qqtqaccatq acccqtqata ccaqcattaq caccqcqtat
                                                                     300
atggaactga gcagcctgcg tagcgaagat acggccgtgt attattgcgc gcgtggtgtt
cattettatt ttgattattg gggccaagge accetggtga eggttagete age
                                                                     353
<210> SEQ ID NO 36
<211> LENGTH: 117
<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 36
Gln Val Gln Leu Val Gln Ser Gly Ala Glu Val Lys Lys Pro Gly Ala
                                   1.0
Ser Val Lys Val Ser Cys Lys Ala Ser Gly Tyr Thr Phe Thr Gly Asn
Ser Met His Trp Val Arg Gln Ala Pro Gly Gln Gly Leu Glu Trp Met
                           40
Gly Thr Ile Phe Pro Tyr Asp Gly Thr Thr Lys Tyr Ala Gln Lys Phe
Gln Gly Arg Val Thr Met Thr Arg Asp Thr Ser Ile Ser Thr Ala Tyr
                   70
Met Glu Leu Ser Ser Leu Arg Ser Glu Asp Thr Ala Val Tyr Tyr Cys
Ala Arg Gly Val His Ser Tyr Phe Asp Tyr Trp Gly Gln Gly Thr Leu
                              105
Val Thr Val Ser Ser
      115
<210> SEQ ID NO 37
<211> LENGTH: 327
<212> TYPE: DNA
<213 > ORGANISM: Homo sapiens
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Thr Ala Arg Ile Ser Cys Ser Gly Asp Asn Leu Arg Gly Tyr Tyr Ala

<400> SEOUENCE: 37 gatatccaga tgacccagag cccgtctagc ctgagcgcga gcgtgggtga tcgtgtgacc 60 attacctgca gagcgagcca gtctattcgt tcttatctgg cttggtacca gcagaaacca 120 ggtaaagcac cgaaactatt aatttataag gcttctaatt tgcaaagcgg ggtcccgtcc 180 cgttttagcg gctctggatc cggcactgat tttaccctga ccattagcag cctgcaacct 240 gaagactttg cggtttatta ttgccatcag tattctgatt ctcctgttac ctttggccag 300 ggtacgaaag ttgaaattaa acgtacg 327 <210> SEQ ID NO 38 <211> LENGTH: 109 <212> TYPE: PRT <213> ORGANISM: Homo sapiens <400> SEQUENCE: 38 Asp Ile Gln Met Thr Gln Ser Pro Ser Ser Leu Ser Ala Ser Val Gly Asp Arg Val Thr Ile Thr Cys Arg Ala Ser Gln Ser Ile Arg Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu Ile Tyr Lys Ala Ser Asn Leu Gln Ser Gly Val Pro Ser Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Gln Pro Glu Asp Phe Ala Val Tyr Tyr Cys His Gln Tyr Ser Asp Ser Pro Val Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr 100 <210> SEQ ID NO 39 <211> LENGTH: 365 <212> TYPE: DNA <213 > ORGANISM: Homo sapiens <400> SEOUENCE: 39 caggtgcaat tgcaacagtc tggtccgggc ctggtgaaac cgagccaaac cctgagcctg 60 acctgtgcga tttccggaga tagcgtgagc tctaattctg ctgcttgggg ttggattcgc 120 cagteteetg ggegtggeet egagtggetg ggeatgatet ateategtag caagtggtat 180 aacgattatg cggtgagcgt gaaaagccgg attaccatca acccggatac ttcgaaaaac 240 cagtttagcc tgcaactgaa cagcgtgacc ccggaagata cggccgtgta ttattgcgcg 300 cgttattctt ctattggtca tatggattat tggggccaag gcaccctggt gacggttagc tcagc 365 <210> SEQ ID NO 40 <211> LENGTH: 121 <212> TYPE: PRT <213 > ORGANISM: Homo sapiens Gln Val Gln Leu Gln Gln Ser Gly Pro Gly Leu Val Lys Pro Ser Gln 10

Thr Leu Ser Leu Thr Cys Ala Ile Ser Gly Asp Ser Val Ser Ser Asn 20 25 30	
Ser Ala Ala Trp Gly Trp Ile Arg Gln Ser Pro Gly Arg Gly Leu Glu 35 40 45	
Trp Leu Gly Met Ile Tyr His Arg Ser Lys Trp Tyr Asn Asp Tyr Ala 50 55 60	
Val Ser Val Lys Ser Arg Ile Thr Ile Asn Pro Asp Thr Ser Lys Asn 65 70 75 80	
Gln Phe Ser Leu Gln Leu Asn Ser Val Thr Pro Glu Asp Thr Ala Val 85 90 95	
Tyr Tyr Cys Ala Arg Tyr Ser Ser Ile Gly His Met Asp Tyr Trp Gly 100 105 110	
Gln Gly Thr Leu Val Thr Val Ser Ser 115 120	
<210> SEQ ID NO 41 <211> LENGTH: 330 <212> TYPE: DNA <213> ORGANISM: Homo sapiens <400> SEQUENCE: 41	
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caggegecag ttettgtgat ttattatgat tetaagegte eetcaggeat eeeggaacge	180
tttageggat ceaacagegg caacacegeg accetgacea ttageggeac teaggeggaa	240
gacgaagcgg attattattg ccaggettat actggtcagt ctattteteg tgtgtttgge	300
ggcggcacga agttaaccgt tcttggccag	330
<210> SEQ ID NO 42 <211> LENGTH: 110 <212> TYPE: PRT <213> ORGANISM: Homo sapiens	
<400> SEQUENCE: 42	
Asp Ile Glu Leu Thr Gln Pro Pro Ser Val Ser Val Ala Pro Gly Gln 1 5 10 15	
Thr Ala Arg Ile Ser Cys Ser Gly Asp Ser Ile Gly Ser Tyr Tyr Ala 20 25 30	
His Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Val Leu Val Ile Tyr 35 40 45	
Tyr Asp Ser Lys Arg Pro Ser Gly Ile Pro Glu Arg Phe Ser Gly Ser 50 60	
Asn Ser Gly Asn Thr Ala Thr Leu Thr Ile Ser Gly Thr Gln Ala Glu 65 70 75 80	
Asp Glu Ala Asp Tyr Tyr Cys Gln Ala Tyr Thr Gly Gln Ser Ile Ser 85 90 95	
Arg Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu Gly Gln 100 105 110	
<210> SEQ ID NO 43 <211> LENGTH: 353 <212> TYPE: DNA <213> ORGANISM: Homo sapiens	

<400> SEQUENCE: 43	
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agetgegegg ceteeggatt tacettttet cettatgtta tgtettgggt gegeeaagee	120
cctgggaagg gtctcgagtg ggtgagctct atctcttctt cttctagcaa tacctattat	180
gcggatagcg tgaaaggccg ttttaccatt tcacgtgata attcgaaaaa caccctgtat	240
ctgcaaatga acagcctgcg tgcggaagat acggccgtgt attattgcgc gcgtggtgat	300
tottatatgt atgatgtttg gggccaaggc accotggtga cggttagctc agc	353
<210> SEQ ID NO 44 <211> LENGTH: 117 <212> TYPE: PRT <213> ORGANISM: Homo sapiens <400> SEQUENCE: 44	
Gln Val Gln Leu Val Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly	
1 5 10 15	
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Pro Tyr 20 25 30	
Val Met Ser Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val 35 40 45	
Ser Ser Ile Ser Ser Ser Ser Ser Asn Thr Tyr Tyr Ala Asp Ser Val	
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr 65 70 75 80	
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys 85 90 95	
Ala Arg Gly Asp Ser Tyr Met Tyr Asp Val Trp Gly Gln Gly Thr Leu 100 105 110	
Val Thr Val Ser Ser 115	
<210> SEQ ID NO 45 <211> LENGTH: 327 <212> TYPE: DNA <213> ORGANISM: Homo sapiens <400> SEQUENCE: 45	
gatatecaga tgacceagag ecegtetage etgagegega gegtgggtga tegtgtgace	60
attacetgea gagegageea ggatattegt aataatetgg ettggtacea geagaaacea	120
ggtaaagcac cgaaactatt aatttatgct gcttcttctt tgcaaagcgg ggtcccgtcc	180
cgttttagcg gctctggatc cggcactgat tttaccctga ccattagcag cctgcaacct	240
gaagactttg cggtttatta ttgccagcag cgtaatggtt ttcctcttac ctttggccag	300
ggtacgaaag ttgaaattaa acgtacg	327
<210> SEQ ID NO 46 <211> LENGTH: 109 <212> TYPE: PRT <213> ORGANISM: Homo sapiens	
<400> SEQUENCE: 46	
Asp Ile Gln Met Thr Gln Ser Pro Ser Ser Leu Ser Ala Ser Val Gly	

1 5 10 15
Asp Arg Val Thr Ile Thr Cys Arg Ala Ser Gln Asp Ile Arg Asn Asn 20 25 30
Leu Ala Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu Ile
35 40 45
Tyr Ala Ala Ser Ser Leu Gln Ser Gly Val Pro Ser Arg Phe Ser Gly 50 60
Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Gln Pro
65 70 75 80
Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Arg Asn Gly Phe Pro Leu 85 90 95
Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr
100 105
<210> SEQ ID NO 47
<211> LENGTH: 365 <212> TYPE: DNA
<213> ORGANISM: Homo sapiens
<400> SEQUENCE: 47
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acctgtgcga tttccggaga tagcgtgagc tctaattctg ctgcttgggg ttggattcgc
cagteteetg ggegtggeet egagtggetg ggeattatet attategtag caagtggtat
aaccattatg cggtgagcgt gaaaagccgg attaccatca acccggatac ttcgaaaaac
cagtttagcc tgcaactgaa cagcgtgacc ccggaagata cggccgtgta ttattgcgcg
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tcagc
<210> SEQ ID NO 48 <211> LENGTH: 121
<212> TYPE: PRT <213> ORGANISM: Homo sapiens
<400> SEQUENCE: 48
Gln Val Gln Leu Gln Gln Ser Gly Pro Gly Leu Val Lys Pro Ser Gln
1 5 10 15
Thr Leu Ser Leu Thr Cys Ala Ile Ser Gly Asp Ser Val Ser Ser Asn 20 25 30
Ser Ala Ala Trp Gly Trp Ile Arg Gln Ser Pro Gly Arg Gly Leu Glu
35 40 45
Trp Leu Gly Ile Ile Tyr Tyr Arg Ser Lys Trp Tyr Asn His Tyr Ala 50 55 60
Val Ser Val Lys Ser Arg Ile Thr Ile Asn Pro Asp Thr Ser Lys Asn
65 70 75 80
Gln Phe Ser Leu Gln Leu Asn Ser Val Thr Pro Glu Asp Thr Ala Val 85 90 95
Tyr Tyr Cys Ala Arg Ser Asn Trp Ser Gly Tyr Phe Asp Tyr Trp Gly
100 105 110
Gln Gly Thr Leu Val Thr Val Ser Ser 115

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<211> LENGTH: 342
<212> TYPE: DNA
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 49
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                                                                     120
taccttcaaa aaccaggtca aagcccgcag ctattaattt atcttggttc taatcgtgcc
                                                                     180
agtggggtcc cggatcgttt tagcggctct ggatccggca ccgattttac cctgaaaatt
                                                                     240
agccgtgtgg aagctgaaga cgtgggcgtg tattattgcc agcagtatga taatgctcct
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attacctttg gccagggtac gaaagttgaa attaaacgta cg
                                                                     342
<210> SEQ ID NO 50
<211> LENGTH: 114
<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 50
Asp Ile Val Met Thr Gln Ser Pro Leu Ser Leu Pro Val Thr Pro Gly
Glu Pro Ala Ser Ile Ser Cys Arg Ser Ser Gln Ser Leu Leu His Ser
Asn Gly Tyr Thr Tyr Leu Ser Trp Tyr Leu Gln Lys Pro Gly Gln Ser
Pro Gln Leu Leu Ile Tyr Leu Gly Ser Asn Arg Ala Ser Gly Val Pro
Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Lys Ile
Ser Arg Val Glu Ala Glu Asp Val Gly Val Tyr Tyr Cys Gln Gln Tyr
Asp Asn Ala Pro Ile Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys
           100
Arg Thr
<210> SEQ ID NO 51
<211> LENGTH: 371
<212> TYPE: DNA
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 51
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acctgtgcga tttccggaga tagcgtgagc tctaattctg ctgcttgggg ttggattcgc
cagteteetg ggegtggeet egagtggetg ggeettatet attategtag caagtggtat
aacgattatg cggtgagcgt gaaaagccgg attaccatca acccggatac ttcgaaaaac
cagtttagcc tgcaactgaa cagcgtgacc ccggaagata cggccgtgta ttattgcgcg
cgttttggtg atactaatcg taatggtact gatgtttggg gccaaggcac cctggtgacg
gttagctcag c
<210> SEQ ID NO 52
<211> LENGTH: 123
<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
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< 400	)> SF	EQUEN	ICE :	52												
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Gln 1	Val	Gln	Leu	Gln 5	Gln	Ser	Gly	Pro	Gly 10	Leu	Val	Lys	Pro	Ser 15	Gln	
Thr	Leu	Ser	Leu 20	Thr	Cys	Ala	Ile	Ser 25	Gly	Asp	Ser	Val	Ser 30	Ser	Asn	
Ser	Ala	Ala 35	Trp	Gly	Trp	Ile	Arg 40	Gln	Ser	Pro	Gly	Arg 45	Gly	Leu	Glu	
Trp	Leu 50	Gly	Leu	Ile	Tyr	Tyr 55	Arg	Ser	ГЛа	Trp	Tyr 60	Asn	Asp	Tyr	Ala	
Val 65	Ser	Val	Lys	Ser	Arg 70	Ile	Thr	Ile	Asn	Pro 75	Asp	Thr	Ser	Lys	Asn 80	
Gln	Phe	Ser	Leu	Gln 85	Leu	Asn	Ser	Val	Thr 90	Pro	Glu	Asp	Thr	Ala 95	Val	
Tyr	Tyr	Cys	Ala 100	Arg	Phe	Gly	Asp	Thr 105	Asn	Arg	Asn	Gly	Thr 110	Asp	Val	
Trp	Gly	Gln 115	Gly	Thr	Leu	Val	Thr 120	Val	Ser	Ser						
		EQ II ENGTH														
		PE: RGANI		Homo	sar	oiens	3									
<400	)> SE	EQUEN	ICE :	53												
gata	tege	cac t	gaco	cago	cc aç	gette	agto	gago	ggct	cac	cago	gtcaç	gag d	catta	accatc	60
tcgt	gtac	gg g	gtact	agca	ag co	gatat	tggt	ggt	tata	att	atgt	gtct	tg (	gtaco	cagcag	120
cato	ccgg	gga a	ggcg	geega	aa ac	ttat	gatt	tat	ggtg	gtta	atta	atcgt	cc o	ctcaç	gcgtg	180
agca	acco	gtt t	tago	ggat	c ca	aaaq	gegge	aac	cacco	gcga	gcct	gaco	cat t	agco	gcctg	240
caaç	ıcgga	aag a	acgaa	agcgg	ga tt	atta	ittgo	tct	tctg	gctg	ataa	agttt	ac t	atgt	ctatt	300
gtgt	ttgg	geg g	gegge	cacga	aa gt	taac	cgtt	ctt	ggcc	ag						339
<211 <212	.> LE :> TY	EQ II ENGTH IPE: RGANI	H: 11 PRT	L3	sar	oiens	3									
< 400	)> SE	EQUEN	ICE :	54												
Asp 1	Ile	Ala	Leu	Thr 5	Gln	Pro	Ala	Ser	Val 10	Ser	Gly	Ser	Pro	Gly 15	Gln	
Ser	Ile	Thr	Ile 20	Ser	Cys	Thr	Gly	Thr 25	Ser	Ser	Asp	Ile	Gly 30	Gly	Tyr	
Asn	Tyr	Val 35	Ser	Trp	Tyr	Gln	Gln 40	His	Pro	Gly	Lys	Ala 45	Pro	Lys	Leu	
Met	Ile 50	Tyr	Gly	Val	Asn	Tyr 55	Arg	Pro	Ser	Gly	Val 60	Ser	Asn	Arg	Phe	
Ser 65	Gly	Ser	Lys	Ser	Gly 70	Asn	Thr	Ala	Ser	Leu 75	Thr	Ile	Ser	Gly	Leu 80	
Gln	Ala	Glu	Asp	Glu 85	Ala	Asp	Tyr	Tyr	ao Gàs	Ser	Ser	Ala	Asp	Lуз 95	Phe	
Thr	Met	Ser	Ile 100	Val	Phe	Gly	Gly	Gly 105	Thr	Lys	Leu	Thr	Val 110	Leu	Gly	

Gln

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<210> SEQ ID NO 55 <211> LENGTH: 306 <212> TYPE: DNA <213 > ORGANISM: Homo sapiens <400> SEOUENCE: 55 gacctgtgcg atttccggag atagcgtgag ctctaattct gctgcttggg gttggattcg 60 ccagtctcct gggcgtggcc tcgagtggct gggcatgatc tattatcgta gcaagtggta 120 taacgattat gcggtgagcg tgaaaagccg gattaccatc aacccggata cttcgaaaaa ccagtttagc ctgcaactga acagcgtgac cccggaagat acggccgtgt attattgcgc gcgtgttaat cagtatactt cttctgatta ttggggccaa ggcaccctgg tgacggttag 300 ctcagc 306 <210> SEQ ID NO 56 <211> LENGTH: 121 <212> TYPE: PRT <213 > ORGANISM: Homo sapiens <400> SEQUENCE: 56 Gln Val Gln Leu Gln Gln Ser Gly Pro Gly Leu Val Lys Pro Ser Gln Thr Leu Ser Leu Thr Cys Ala Ile Ser Gly Asp Ser Val Ser Ser Asn Ser Ala Ala Trp Gly Trp Ile Arg Gln Ser Pro Gly Arg Gly Leu Glu 35 40 45 Trp Leu Gly Met Ile Tyr Tyr Arg Ser Lys Trp Tyr As<br/>n Asp Tyr Ala 50  $\,$  60 Val Ser Val Lys Ser Arg Ile Thr Ile Asn Pro Asp Thr Ser Lys Asn 65 70 75 80 Gln Phe Ser Leu Gln Leu Asn Ser Val Thr Pro Glu Asp Thr Ala Val 90 Tyr Tyr Cys Ala Arg Val Asn Gln Tyr Thr Ser Ser Asp Tyr Trp Gly 100 Gln Gly Thr Leu Val Thr Val Ser Ser 115 <210> SEQ ID NO 57 <211> LENGTH: 327 <212> TYPE: DNA <213 > ORGANISM: Homo sapiens <400> SEQUENCE: 57 gatatccaga tgacccagag cccgtctagc ctgagcgcga gcgtgggtga tcgtgtgacc attacctgca gagcgagcca gcctatttat aattctctgt cttggtacca gcagaaacca ggtaaagcac cgaaactatt aatttatggt gtttctaatt tgcaaagcgg ggtcccgtcc cgttttagcg gctctggatc cggcactgat tttaccctga ccattagcag cctgcaacct gaagactttg cggtttatta ttgccttcag gttgataatc ttcctattac ctttggccag ggtacgaaag ttgaaattaa acgtacg

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<211> LENGTH: 109
<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 58
Asp Ile Gln Met Thr Gln Ser Pro Ser Ser Leu Ser Ala Ser Val Gly
                                   10
Asp Arg Val Thr Ile Thr Cys Arg Ala Ser Gln Pro Ile Tyr Asn Ser
                              25
Leu Ser Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu Ile
Tyr Gly Val Ser Asn Leu Gln Ser Gly Val Pro Ser Arg Phe Ser Gly
Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Gln Pro
Glu Asp Phe Ala Val Tyr Tyr Cys Leu Gln Val Asp Asn Leu Pro Ile
Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr
<210> SEQ ID NO 59
<211> LENGTH: 374
<212> TYPE: DNA
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 59
caggtgcaat tgcaacagtc tggtccgggc ctggtgaaac cgagccaaac cctgagcctg
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acctgtgcga tttccggaga tagcgtgagc tctaattctg ctgcttggtc ttggattcgc
                                                                     120
                                                                     180
cagtetectg ggegtggeet egagtggetg ggeatgatet tittategtag caagtggaat
aacgattatg cggtgagcgt gaaaagccgg attaccatca acccggatac ttcgaaaaac
                                                                     240
cagtttagcc tgcaactgaa cagcgtgacc ccggaagata cggccgtgta ttattgcgcg
                                                                     300
cgtgttaatg ctaatggtta ttatgcttat gttgatcttt ggggccaagg caccctggtg
                                                                     360
acggttagct cagc
                                                                     374
<210> SEQ ID NO 60
<211> LENGTH: 124
<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 60
Gln Val Gln Leu Gln Gln Ser Gly Pro Gly Leu Val Lys Pro Ser Gln
                                   10
Thr Leu Ser Leu Thr Cys Ala Ile Ser Gly Asp Ser Val Ser Ser Asn
Ser Ala Ala Trp Ser Trp Ile Arg Gln Ser Pro Gly Arg Gly Leu Glu
Trp Leu Gly Met Ile Phe Tyr Arg Ser Lys Trp Asn Asn Asp Tyr Ala
Val Ser Val Lys Ser Arg Ile Thr Ile Asn Pro Asp Thr Ser Lys Asn
Gln Phe Ser Leu Gln Leu Asn Ser Val Thr Pro Glu Asp Thr Ala Val
Tyr Tyr Cys Ala Arg Val Asn Ala Asn Gly Tyr Tyr Ala Tyr Val Asp
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100	105 110	
Leu Trp Gly Gln Gly Thr Leu Val 115 120	Thr Val Ser Ser	
<210> SEQ ID NO 61 <211> LENGTH: 330 <212> TYPE: DNA <213> ORGANISM: Homo sapiens		
<400> SEQUENCE: 61		
gatatogtgo tgacccagag cooggogacc	c ctgagcctgt ctccgggcga acgtgcgacc	60
ctgagctgca gagcgagcca gtctgtttct	teteagtate tggettggta eeageagaaa	120
ccaggtcaag caccgcgtct attaatttat	getgettett etegtgeaae tggggteeeg	180
gegegtttta geggetetgg ateeggeacg	g gattttaccc tgaccattag cagcctggaa	240
cctgaagact ttgcggttta ttattgccag	g caggatteta atetteetge tacetttgge	300
cagggtacga aagttgaaat taaacgtacg	3	330
<210> SEQ ID NO 62 <211> LENGTH: 110 <212> TYPE: PRT <213> ORGANISM: Homo sapiens		
<400> SEQUENCE: 62		
Asp Ile Val Leu Thr Gln Ser Pro 1 5	Ala Thr Leu Ser Leu Ser Pro Gly 10 15	
Glu Arg Ala Thr Leu Ser Cys Arg . 20	Ala Ser Gln Ser Val Ser Ser Gln 25 30	
Tyr Leu Ala Trp Tyr Gln Gln Lys 35 40	Pro Gly Gln Ala Pro Arg Leu Leu 45	
Ile Tyr Ala Ala Ser Ser Arg Ala 50 55	Thr Gly Val Pro Ala Arg Phe Ser	
Gly Ser Gly Ser Gly Thr Asp Phe 65 70	Thr Leu Thr Ile Ser Ser Leu Glu 75 80	
Pro Glu Asp Phe Ala Val Tyr Tyr 85	Cys Gln Gln Asp Ser Asn Leu Pro 90 95	
Ala Thr Phe Gly Gln Gly Thr Lys	Val Glu Ile Lys Arg Thr 105 110	
<210> SEQ ID NO 63 <211> LENGTH: 351 <212> TYPE: DNA <213> ORGANISM: Homo sapiens		
<400> SEQUENCE: 63		
caggtgcaat tggtggaaag cggcggcggc	c ctggtgcaac cgggcggcag cctgcgtctg	60
agetgegegg ceteeggatt tacettttat	: aagtatgcta tgcattgggt gcgccaagcc	120
cctgggaagg gtctcgagtg ggtgagcggt	atccagtatg atggtagcta tacctattat	180
gcggatagcg tgaaaggccg ttttaccatt	tcacgtgata attcgaaaaa caccctgtat	240
ctgcaaatga acagcctgcg tgcggaagat	acggccgtgt attattgcgc gcgttattat	300
tgtaagtgtg ttgatctttg gggccaaggc	c accetggtga eggttagete a	351

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<211> LENGTH: 117
<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 64
Gln Val Gln Leu Val Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
                                   10
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Tyr Lys Tyr
                              25
Ala Met His Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
                           40
Ser Gly Ile Gln Tyr Asp Gly Ser Tyr Thr Tyr Tyr Ala Asp Ser Val
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys
Ala Arg Tyr Tyr Cys Lys Cys Val Asp Leu Trp Gly Gln Gly Thr Leu
Val Thr Val Ser Ser
<210> SEQ ID NO 65
<211> LENGTH: 327
<212> TYPE: DNA
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 65
gatategaac tgacceagee geetteagtg agegttgeac caggteagae egegegtate
tcgtgtagcg gcgataatat tcgtaagttt tatgttcatt ggtaccagca gaaacccggg
caggegeeag ttettgtgat ttatggtaet aataagegte ceteaggeat eeeggaaege
                                                                     180
tttageggat ccaacagegg caacacegeg accetgacea ttageggeac tcaggeggaa
                                                                     240
gacgaagcgg attattattg ccagtcttat gattctaagt ttaatactgt gtttggcggc
                                                                     300
                                                                     327
ggcacgaagt taaccgttct tggccag
<210> SEQ ID NO 66
<211> LENGTH: 109
<212> TYPE: PRT
<213> ORGANISM: Homo sapiens
<400> SEQUENCE: 66
Asp Ile Glu Leu Thr Gln Pro Pro Ser Val Ser Val Ala Pro Gly Gln
Thr Ala Arg Ile Ser Cys Ser Gly Asp Asn Ile Arg Lys Phe Tyr Val
His Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Val Leu Val Ile Tyr
Gly Thr Asn Lys Arg Pro Ser Gly Ile Pro Glu Arg Phe Ser Gly Ser
Asn Ser Gly Asn Thr Ala Thr Leu Thr Ile Ser Gly Thr Gln Ala Glu
Asp Glu Ala Asp Tyr Tyr Cys Gln Ser Tyr Asp Ser Lys Phe Asn Thr
```

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Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu Gly Gln
           100
                               105
<210> SEQ ID NO 67
<211> LENGTH: 359
<212> TYPE: DNA
<213> ORGANISM: Homo sapiens
<400> SEOUENCE: 67
caggtgcaat tggtggaaag cggcggcggc ctggtgcaac cgggcggcag cctgcgtctg
                                                                      60
agetgegegg ceteeggatt tacettttet tettatgeta tgaattgggt gegeeaagee
                                                                     120
cctgggaagg gtctcgagtg ggtgagcgct atcctttctg atggtagctc tacctcttat
geggatageg tgaaaggeeg ttttaccatt teaegtgata attegaaaaa caccetgtat
                                                                     240
ctgcaaatga acagcctgcg tgcggaagat acggccgtgt attattgcgc gcgttatcct
gattggggtt ggtatactga tgtttggggc caaggcaccc tggtgacggt tagctcagc
<210> SEQ ID NO 68
<211> LENGTH: 119
<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 68
Gln Val Gln Leu Val Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Ser Tyr
                               25
Ala Met Asn Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
                          40
Ser Ala Ile Leu Ser Asp Gly Ser Ser Thr Ser Tyr Ala Asp Ser Val
                       55
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr
                    70
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys
Ala Arg Tyr Pro Asp Trp Gly Trp Tyr Thr Asp Val Trp Gly Gln Gly
           100
                               105
Thr Leu Val Thr Val Ser Ser
       115
<210> SEQ ID NO 69
<211> LENGTH: 330
<212> TYPE: DNA
<213 > ORGANISM: Homo sapiens
<400> SEOUENCE: 69
gatategaac tgacceagec geetteagtg agegttgeac caggteagac egegegtate
tcgtgtagcg gcgatgctct tcgtaagcat tatgtttatt ggtaccagca gaaacccggg
caggegecag ttettgtgat ttatggtgat aataategte ceteaggeat eeeggaaege
tttageggat ccaacagegg caacacegeg accetgacea ttageggeac tcaggeggaa
gacgaagcgg attattattg ccagtcttat gataagcctt atcctattct tgtgtttggc
ggcggcacga agttaaccgt tcttggccag
```

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<210> SEQ ID NO 70
<211> LENGTH: 110
<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 70
Asp Ile Glu Leu Thr Gln Pro Pro Ser Val Ser Val Ala Pro Gly Gln
                                   1.0
Thr Ala Arg Ile Ser Cys Ser Gly Asp Ala Leu Arg Lys His Tyr Val
Tyr Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Val Leu Val Ile Tyr
Gly Asp Asn Asn Arg Pro Ser Gly Ile Pro Glu Arg Phe Ser Gly Ser
Asn Ser Gly Asn Thr Ala Thr Leu Thr Ile Ser Gly Thr Gln Ala Glu
Asp Glu Ala Asp Tyr Tyr Cys Gln Ser Tyr Asp Lys Pro Tyr Pro Ile
Leu Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu Gly Gln
<210> SEQ ID NO 71
<211> LENGTH: 356
<212> TYPE: DNA
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 71
caggtgcaat tggtggaaag cggcggcggc ctggtgcaac cgggcggcag cctgcgtctg
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agetgegegg ceteeggatt tacettttet tettatgeta tgaettgggt gegeeaagee
cctgggaagg gtctcgagtg ggtgagcaat atctcttatt ctggtagcaa tacctattat
geggatageg tgaaaggeeg ttttaccatt teaegtgata attegaaaaa caccetgtat
                                                                     240
ctgcaaatga acagcctgcg tgcggaagat acggccgtgt attattgcgc gcgtgttggt
                                                                     300
tattattatg gttttgatta ttggggccaa ggcaccctgg tgacggttag ctcagc
                                                                     356
<210> SEQ ID NO 72
<211> LENGTH: 118
<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
<400> SEOUENCE: 72
Gln Val Gln Leu Val Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Ser Tyr
                               25
Ala Met Thr Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
Ser Asn Ile Ser Tyr Ser Gly Ser Asn Thr Tyr Tyr Ala Asp Ser Val
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys
Ala Arg Val Gly Tyr Tyr Gly Phe Asp Tyr Trp Gly Gln Gly Thr
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Leu Val Thr Val Ser Ser 115 <210> SEO ID NO 73 <211> LENGTH: 330 <212> TYPE: DNA <213 > ORGANISM: Homo sapiens <400> SEQUENCE: 73 gatatcgtgc tgacccagag cccggcgacc ctgagcctgt ctccgggcga acgtgcgacc 60 ctgagetgea gagegageea gaatgtttet tetaattate tggettggta eeageagaaa 120 ccaggtcaag caccgcgtct attaatttat gatgcttcta atcgtgcaac tggggtcccg gegegtttta geggetetgg atceggeacg gattttacce tgaccattag cageetggaa cctgaagact ttgcggttta ttattgccag cagttttatg attctcctca gacctttggc cagggtacga aagttgaaat taaacgtacg 330 <210> SEQ ID NO 74 <211> LENGTH: 110 <212> TYPE: PRT <213 > ORGANISM: Homo sapiens <400> SEQUENCE: 74 Asp Ile Val Leu Thr Gln Ser Pro Ala Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Asn Val Ser Ser Asn Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Asp Ala Ser Asn Arg Ala Thr Gly Val Pro Ala Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Phe Tyr Asp Ser Pro Gln Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr 100 <210> SEO ID NO 75 <211> LENGTH: 365 <212> TYPE: DNA <213> ORGANISM: Homo sapiens <400> SEOUENCE: 75 caggtgcaat tgcaacagtc tggtccgggc ctggtgaaac cgagccaaac cctgagcctg 60 acctgtgcga tttccggaga tagcgtgagc tctaattctg ctgcttggtc ttggattcgc cagtetectg ggegtggeet egagtggetg ggetttatet attategtag caagtggtat aacgattatg cggtgagcgt gaaaagccgg attaccatca acccggatac ttcgaaaaac cagtttagcc tgcaactgaa cagcgtgacc ccggaagata cggccgtgta ttattgcgcg cgtcataatc ctgatcttgg ttttgattat tggggccaag gcaccctggt gacggttagc tcagc 365

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<210> SEQ ID NO 76
<211> LENGTH: 121
<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 76
Gln Val Gln Leu Gln Gln Ser Gly Pro Gly Leu Val Lys Pro Ser Gln
Thr Leu Ser Leu Thr Cys Ala Ile Ser Gly Asp Ser Val Ser Ser Asn
Ser Ala Ala Trp Ser Trp Ile Arg Gln Ser Pro Gly Arg Gly Leu Glu
Trp Leu Gly Phe Ile Tyr Tyr Arg Ser Lys Trp Tyr Asn Asp Tyr Ala
Val Ser Val Lys Ser Arg Ile Thr Ile Asn Pro Asp Thr Ser Lys Asn
Gln Phe Ser Leu Gln Leu Asn Ser Val Thr Pro Glu Asp Thr Ala Val
Tyr Tyr Cys Ala Arg His Asn Pro Asp Leu Gly Phe Asp Tyr Trp Gly
Gln Gly Thr Leu Val Thr Val Ser Ser
<210> SEQ ID NO 77
<211> LENGTH: 327
<212> TYPE: DNA
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 77
qatateqtqc tqacccaqaq cccqqcqacc ctqaqcctqt ctccqqqcqa acqtqcqacc
                                                                      60
ctqaqctqca qaqcqaqcca qtatqttact tcttcttatc tqqcttqqta ccaqcaqaaa
                                                                     120
ccaggtcaag caccgcgtct attaatttat ggttcttctc gtgcaactgg ggtcccggcg
                                                                     180
cqttttaqcq qctctqqatc cqqcacqqat tttaccctqa ccattaqcaq cctqqaacct
                                                                     240
gaagactttg cgacttatta ttgccagcag tattcttctt ctcctattac ctttggccag
                                                                     300
ggtacgaaag ttgaaattaa acgtacg
                                                                     327
<210> SEQ ID NO 78
<211> LENGTH: 109
<212> TYPE: PRT
<213> ORGANISM: Homo sapiens
<400> SEQUENCE: 78
Asp Ile Val Leu Thr Gln Ser Pro Ala Thr Leu Ser Leu Ser Pro Gly
                                   10
Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Tyr Val Thr Ser Ser
Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu
Ile Tyr Gly Ser Ser Arg Ala Thr Gly Val Pro Ala Arg Phe Ser Gly
Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Glu Pro
Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Tyr Ser Ser Pro Ile
```

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Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr
           100
<210> SEQ ID NO 79
<211> LENGTH: 362
<212> TYPE: DNA
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 79
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acctgtgcga tttccggaga tagcgtgagc tcttcttctg ctgcttggtc ttggattcgc
                                                                     120
cagtetectg ggegtggeet egagtggetg ggeattatet attategtag caagtggtat
aacgattatg cggtgagcgt gaaaagccgg attaccatca acccggatac ttcgaaaaac
cagtttagcc tgcaactgaa cagcgtgacc ccggaagata cggccgtgta ttattgcgcg
cgtcattcta tggttggttt tgatgtttgg ggccaaggca ccctggtgac ggttagctca
<210> SEQ ID NO 80
<211> LENGTH: 120
<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 80
Gln Val Gln Leu Gln Gln Ser Gly Pro Gly Leu Val Lys Pro Ser Gln
                                  10
Thr Leu Ser Leu Thr Cys Ala Ile Ser Gly Asp Ser Val Ser Ser Ser
                               25
Ser Ala Ala Trp Ser Trp Ile Arg Gln Ser Pro Gly Arg Gly Leu Glu
Trp Leu Gly Ile Ile Tyr Tyr Arg Ser Lys Trp Tyr Asn Asp Tyr Ala
Val Ser Val Lys Ser Arg Ile Thr Ile Asn Pro Asp Thr Ser Lys Asn
Gln Phe Ser Leu Gln Leu Asn Ser Val Thr Pro Glu Asp Thr Ala Val
                                    90
Tyr Tyr Cys Ala Arg His Ser Met Val Gly Phe Asp Val Trp Gly Gln \,
           100
                               105
Gly Thr Leu Val Thr Val Ser Ser
       115
<210> SEQ ID NO 81
<211> LENGTH: 330
<212> TYPE: DNA
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 81
gatatcgaac tgacccagcc gccttcagtg agcgttgcac caggtcagac cgcgcgtatc
tcgtgtagcg gcgataatct tggtacttat tatgttcatt ggtaccagca gaaacccggg
caggegecag ttettgtgat ttatggtgat aataategte ceteaggeat eeeggaacge
tttageggat ccaacagegg caacacegeg accetgacea ttageggeac tcaggeggaa
gacgaagcgg attattattg ccagacttat gattctaata atgagtctat tgtgtttggc
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ggcggcacga agttaaccgt tcttggccag	330
<210> SEQ ID NO 82 <211> LENGTH: 110 <212> TYPE: PRT <213> ORGANISM: Homo sapiens	
<400> SEQUENCE: 82	
Asp Ile Glu Leu Thr Gln Pro Pro Ser Val Ser Val Ala Pro Gly Gln 1 5 10 15	
Thr Ala Arg Ile Ser Cys Ser Gly Asp Asn Leu Gly Thr Tyr Tyr Val	
His Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Val Leu Val Ile Tyr 35 40 45	
Gly Asp Asn Asn Arg Pro Ser Gly Ile Pro Glu Arg Phe Ser Gly Ser 50 60	
Asn Ser Gly Asn Thr Ala Thr Leu Thr Ile Ser Gly Thr Gln Ala Glu 65 70 75 80	
Asp Glu Ala Asp Tyr Tyr Cys Gln Thr Tyr Asp Ser Asn Asn Glu Ser 85 90 95	
Ile Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu Gly Gln 100 105 110	
<210> SEQ ID NO 83 <211> LENGTH: 368 <212> TYPE: DNA <213> ORGANISM: Homo sapiens	
<400> SEQUENCE: 83	
caggtgcaat tggtggaaag cggcggcggc ctggtgcaac cgggcggcag cctgcgtctg	60
agetgegegg ceteeggatt taeetttaat tettatgeta tgtettgggt gegeeaagee	120
cctgggaagg gtctcgagtg ggtgagcaat atctcttcta attctagcaa tacctattat	180
geggatageg tgaaaggeeg ttttaccatt teaegtgata attegaaaaa caccetgtat	240
ctgcaaatga acagcctgcg tgcggaagat acggccgtgt attattgcgc gcgtaagggt	300
ggtggtgagc atggtttttt tccttctgat atttggggcc aaggcaccct ggtgacggtt	360
ageteage	368
<210> SEQ ID NO 84 <211> LENGTH: 122 <212> TYPE: PRT <213> ORGANISM: Homo sapiens	
<400> SEQUENCE: 84	
Gln Val Gln Leu Val Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly 1 10 15	
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Asn Ser Tyr 20 25 30	
Ala Met Ser Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val 35 40 45	
Ser Asn Ile Ser Ser Asn Ser Ser Asn Thr Tyr Tyr Ala Asp Ser Val 50 55 60	
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr 65 70 75 80	

Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys 85 90 95	
Ala Arg Lys Gly Gly Gly Glu His Gly Phe Phe Pro Ser Asp Ile Trp 100 105 110	
Gly Gln Gly Thr Leu Val Thr Val Ser Ser 115 120	
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togtgtacgg gtactagcag cgatcttggt ggttttaata ctgtgtcttg gtaccagcag	120
cateceggga aggegeegaa aettatgatt tattetgttt ettetegtee eteaggegtg	180
agcaaccgtt ttagcggatc caaaagcggc aacaccgcga gcctgaccat tagcggcctg	240
caagcggaag acgaagcgga ttattattgc cagtettatg atettaataa tettgtgttt	300
ggcggcggca cgaagttaac cgttcttggc cag	333
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Ser Ile Thr Ile Ser Cys Thr Gly Thr Ser Ser Asp Leu Gly Gly Phe 20 25 30	
Asn Thr Val Ser Trp Tyr Gln Gln His Pro Gly Lys Ala Pro Lys Leu 35 40 45	
Met Ile Tyr Ser Val Ser Ser Arg Pro Ser Gly Val Ser Asn Arg Phe 50 55 60	
Ser Gly Ser Lys Ser Gly Asn Thr Ala Ser Leu Thr Ile Ser Gly Leu 65 70 75 80	
Gln Ala Glu Asp Glu Ala Asp Tyr Tyr Cys Gln Ser Tyr Asp Leu Asn 85 90 95	
Asn Leu Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu Gly Gln 100 105 110	
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cctgggaagg gtctcgagtg ggtgagcgct atcaagtctg atggtagcaa tacctattat	180
goggatagog tgaaaggoog ttttaccatt toacgtgata attogaaaaa caccotgtat	240
ctgcaaatga acagcctgcg tgcggaagat acggccgtgt attattgcgc gcgtaatgat	300

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tetggttggt ttgatgtttg gggeeaagge accetggtga eggttagete age	353
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Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Asn Ser Tyr 20 25 30	
Ala Met Thr Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val 35 40 45	
Ser Ala Ile Lys Ser Asp Gly Ser Asn Thr Tyr Tyr Ala Asp Ser Val 50 55 60	
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr 65 70 75 80	
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys 85 90 95	
Ala Arg Asn Asp Ser Gly Trp Phe Asp Val Trp Gly Gln Gly Thr Leu 100 105 110	
Val Thr Val Ser Ser 115	
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ctgagctgca gagcgagcca gtctgtttct tctttttatc tggcttggta ccagcagaaa	120
ccaggtcaag caccgcgtct attaatttat ggttcttctt ctcgtgcaac tggggtcccg	180
gcgcgtttta gcggctctgg atccggcacg gattttaccc tgaccattag cagcctggaa	240
cctgaagact ttgcgactta ttattgccag cagtatgatt ctactccttc tacctttggc	300
cagggtacga aagttgaaat taaacgtacg	330
<210> SEQ ID NO 90 <211> LENGTH: 110 <212> TYPE: PRT <213> ORGANISM: Homo sapiens	
<400> SEQUENCE: 90	
Asp Ile Val Leu Thr Gln Ser Pro Ala Thr Leu Ser Leu Ser Pro Gly 1 10 15	
Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Ser Ser Phe 20 25 30	
Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu 35 40 45	
Ile Tyr Gly Ser Ser Ser Arg Ala Thr Gly Val Pro Ala Arg Phe Ser 50 60	
Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Glu 65 70 70 80	

Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Tyr Asp Ser Thr Pro Ser Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr 100 105 <210> SEQ ID NO 91 <211> LENGTH: 365 <212> TYPE: DNA <213> ORGANISM: Homo sapiens <400> SEQUENCE: 91 caggtgcaat tgcaacagtc tggtccgggc ctggtgaaac cgagccaaac cctgagcctg 60 acctgtgcga tttccggaga tagcgtgagc tctaatggtg ctgcttgggg ttggattcgc cagteteetg ggegtggeet egagtggetg ggetttatet ategtegtag caagtggtat aactcttatg cggtgagcgt gaaaagccgg attaccatca acccggatac ttcgaaaaac cagtttagcc tgcaactgaa cagcgtgacc ccggaagata cggccgtgta ttattgcgcg cgtcaggatg gtatgggtgg tatggattet tggggccaag gcaccetggt gaeggttage tcagc <210> SEQ ID NO 92 <211> LENGTH: 121 <212> TYPE: PRT <213 > ORGANISM: Homo sapiens <400> SEQUENCE: 92 Gln Val Gln Leu Gln Gln Ser Gly Pro Gly Leu Val Lys Pro Ser Gln Thr Leu Ser Leu Thr Cys Ala Ile Ser Gly Asp Ser Val Ser Ser Asn Gly Ala Ala Trp Gly Trp Ile Arg Gln Ser Pro Gly Arg Gly Leu Glu 40 Trp Leu Gly Phe Ile Tyr Arg Arg Ser Lys Trp Tyr Asn Ser Tyr Ala Val Ser Val Lys Ser Arg Ile Thr Ile Asn Pro Asp Thr Ser Lys Asn Gln Phe Ser Leu Gln Leu Asn Ser Val Thr Pro Glu Asp Thr Ala Val Tyr Tyr Cys Ala Arg Gln Asp Gly Met Gly Gly Met Asp Ser Trp Gly 100 105 Gln Gly Thr Leu Val Thr Val Ser Ser 115 <210> SEQ ID NO 93 <211> LENGTH: 327 <212> TYPE: DNA <213 > ORGANISM: Homo sapiens <400> SEQUENCE: 93 gatategaac tgacceagee geetteagtg agegttgeac caggteagac egegegtate tcgtgtagcg gcgataatat tggttctcgt tatgcttatt ggtaccagca gaaacccggg caggegecag ttgttgtgat ttatgatgat tetgategte ceteaggeat eeeggaacge tttaqcqqat ccaacaqcqq caacaccqcq accctqacca ttaqcqqcac tcaqqcqqaa

gacgaagcgg attattattg cgctgcttat actttttatg ctcgtactgt gtttggcggc	300
ggcacgaagt taaccgttct tggccag	327
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Thr Ala Arg Ile Ser Cys Ser Gly Asp Asn Ile Gly Ser Arg Tyr Ala 20 25 30	
Tyr Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Val Val Val Ile Tyr 35 40 45	
Asp Asp Ser Asp Arg Pro Ser Gly Ile Pro Glu Arg Phe Ser Gly Ser 50 55 60	
Asn Ser Gly Asn Thr Ala Thr Leu Thr Ile Ser Gly Thr Gln Ala Glu 65 70 75 80	
Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Tyr Thr Phe Tyr Ala Arg Thr 85 90 95	
Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu Gly Gln 100 105	
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agctgcgcgg cctccggatt taccttttct aattattatc tttcttgggt gcgccaagcc	120
cctgggaagg gtctcgagtg ggtgagcggt atctcttata atggtagctc taccaattat	180
gcggatagcg tgaaaggccg ttttaccatt tcacgtgata attcgaaaaa caccctgtat	240
ctgcaaatga acagcctgcg tgcggaagat acggccgtgt attattgcgc gcgtatgtgg	300
cgttattctc ttggtgctga ttcttggggc caaggcaccc tggtgacggt tagctcagc	359
<210> SEQ ID NO 96 <211> LENGTH: 119 <212> TYPE: PRT <213> ORGANISM: Homo sapiens	
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Gln Val Gln Leu Val Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly 1 5 10 15	
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Asn Tyr 20 25 30	
Tyr Leu Ser Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val 35 40 45	
Ser Gly Ile Ser Tyr Asn Gly Ser Ser Thr Asn Tyr Ala Asp Ser Val 50 55 60	
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr 65 70 75 80	

Ala Arg Met Trp Arg Tyr Ser Leu Gly Ala Asp Ser Trp Gly Gln Gly 100 105 Thr Leu Val Thr Val Ser Ser 115 <210> SEQ ID NO 97 <211> LENGTH: 327 <212> TYPE: DNA <213 > ORGANISM: Homo sapiens <400> SEQUENCE: 97 gatategaac tgacceagee geetteagtg agegttgeac caggteagac egegegtate tcgtgtagcg gcgataatat tggttctaag tatgttcatt ggtaccagca gaaacccggg caggegecag tigtigtgat tiatgaggat tetgategic ceteaggeat eeeggaacge tttageggat ccaacagegg caacacegeg accetgacea ttageggeac tcaggeggaa gacgaagcgg attattattg ccagtcttgg gataagtctg agggttatgt gtttggcggc ggcacgaagt taaccgttct tggccag <210> SEQ ID NO 98 <211> LENGTH: 109 <212> TYPE: PRT <213 > ORGANISM: Homo sapiens <400> SEQUENCE: 98 Asp Ile Glu Leu Thr Gln Pro Pro Ser Val Ser Val Ala Pro Gly Gln Thr Ala Arg Ile Ser Cys Ser Gly Asp Asn Ile Gly Ser Lys Tyr Val $20 \hspace{1.5cm} 25 \hspace{1.5cm} 30$ His Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Val Val Val Ile Tyr Glu Asp Ser Asp Arg Pro Ser Gly Ile Pro Glu Arg Phe Ser Gly Ser Asn Ser Gly Asn Thr Ala Thr Leu Thr Ile Ser Gly Thr Gln Ala Glu Asp Glu Ala Asp Tyr Tyr Cys Gln Ser Trp Asp Lys Ser Glu Gly Tyr Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu Gly Gln 100 <210> SEQ ID NO 99 <211> LENGTH: 371 <212> TYPE: DNA <213 > ORGANISM: Homo sapiens <400> SEQUENCE: 99 caggtgcaat tggtggaaag cggcggcggc ctggtgcaac cgggcggcag cctgcgtctg agctgcgcgg cctccggatt tacctttaat aataatgcta tttcttgggt gcgccaagcc cctgggaagg gtctcgagtg ggtgagcgct atcaattctt cttctagctc tacctcttat qcqqataqcq tqaaaqqccq ttttaccatt tcacqtqata attcqaaaaa caccctqtat ctgcaaatga acagcctgcg tgcggaagat acggccgtgt attattgcgc gcgtggtcat

Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys

categtggte attettggge ttettttatt gattattggg gecaaggeae eetggtgaeg	360
gttagctcag c	371
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Gln Val Gln Leu Val Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly 1 5 10 15	
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Asn Asn Asn 20 25 30	
Ala Ile Ser Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val	
Ser Ala Ile Asn Ser Ser Ser Ser Ser Thr Ser Tyr Ala Asp Ser Val	
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr 65 70 75 80	
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys 85 90 95	
Ala Arg Gly His His Arg Gly His Ser Trp Ala Ser Phe Ile Asp Tyr	
Trp Gly Gln Gly Thr Leu Val Thr Val Ser Ser	
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caggcgccag ttcttgtgat ttattctaag tctgagcgtc cctcaggcat cccggaacgc	180
tttageggat ecaacagegg caacacegeg accetgacea ttageggeae teaggeggaa	240
gacgaagcgg attattattg ctcttcttat actcttaatc ctaatcttaa ttatgtgttt	300
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Thr Ala Arg Ile Ser Cys Ser Gly Asp Asn Leu Arg Asp Lys Tyr Ala 20 25 30	
Ser Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Val Leu Val Ile Tyr 35 40 45	
Ser Lys Ser Glu Arg Pro Ser Gly Ile Pro Glu Arg Phe Ser Gly Ser 50 55 60	

Asn Ser Gly Asn Thr Ala Thr Leu Thr Ile Ser Gly Thr Gln Ala Glu Asp Glu Ala Asp Tyr Tyr Cys Ser Ser Tyr Thr Leu Asn Pro Asn Leu Asn Tyr Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu Gly Gln <210> SEQ ID NO 103 <211> LENGTH: 347 <212> TYPE: DNA <213 > ORGANISM: Homo sapiens <400> SEQUENCE: 103 caggtgcaat tggtggaaag cggcggcggc ctggtgcaac cgggcggcag cctgcgtctg agetgegegg ceteeggatt tacettttet tettattgga tgeattgggt gegeeaagee cctgggaagg gtctcgagtg ggtgagctct atctcttatg attctagcaa tacctattat gcggatagcg tgaaaggccg ttttaccatt tcacgtgata attcgaaaaa caccctgtat ctgcaaatga acagcctgcg tgcggaagat acggccgtgt attattgcgc gcgttatggt ggtatggatt attggggcca aggcaccctg gtgacggtta gctcagc <210> SEQ ID NO 104 <211> LENGTH: 115 <212> TYPE: PRT <213 > ORGANISM: Homo sapiens <400> SEQUENCE: 104 Gln Val Gln Leu Val Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly 10 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Ser Tyr Trp Met His Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val Ser Ser Ile Ser Tyr Asp Ser Ser Asn Thr Tyr Tyr Ala Asp Ser Val Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys Ala Arg Tyr Gly Gly Met Asp Tyr Trp Gly Gln Gly Thr Leu Val Thr Val Ser Ser 115 <210> SEQ ID NO 105 <211> LENGTH: 327 <212> TYPE: DNA <213 > ORGANISM: Homo sapiens <400> SEQUENCE: 105 gatategaac tgacceagec ggetteagtg agegttgeac caggteagac egegegtate tcgtgtagcg gcgataatct tcgttctaag tatgctcatt ggtaccagca gaaacccggg caqqqqcaq ttcttqtqat ttatqqtqat aataatcqtc cctcaqqcat cccqqaacqc

tttageggat ccaacagegg caacacegeg accetgacca ttageggeac tcaggeggaa 240
gacgaagegg attattattg etetgettat getatgggtt etteteetgt gtttggegge 300
ggcacgaagt taaccgttct tggccag 327
<210> SEQ ID NO 106 <211> LENGTH: 109 <212> TYPE: PRT <213> ORGANISM: Homo sapiens
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Thr Ala Arg Ile Ser Cys Ser Gly Asp Asn Leu Arg Ser Lys Tyr Ala 20 25 30
His Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Val Leu Val Ile Tyr 35 40 45
Gly Asp Asn Asn Arg Pro Ser Gly Ile Pro Glu Arg Phe Ser Gly Ser 50 60
Asn Ser Gly Asn Thr Ala Thr Leu Thr Ile Ser Gly Thr Gln Ala Glu 65 70 75 80
Asp Glu Ala Asp Tyr Tyr Cys Ser Ala Tyr Ala Met Gly Ser Ser Pro 85 90 95
Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu Gly Gln 100 105
<210> SEQ ID NO 107 <211> LENGTH: 356 <212> TYPE: DNA <213> ORGANISM: Homo sapiens
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gcggatagcg tgaaaggccg ttttaccatt tcacgtgata attcgaaaaa caccctgtat 240
ctgcaaatga acagcctgcg tgcggaagat acggccgtgt attattgcgc gcgtggtctt 300
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Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Ser Tyr 20 25 30
Gly Met His Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val 35 40 45
Ser Asn Ile Ser Tyr Met Gly Ser Asn Thr Asn Tyr Ala Asp Ser Val 50 55 60
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr

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65     70     75     80
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys 85 90 95
Ala Arg Gly Leu Phe Pro Gly Tyr Phe Asp Tyr Trp Gly Gln Gly Thr 100 105 110
Leu Val Thr Val Ser Ser 115
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ggtaaagcac cgaaactatt aatttatggt acttettett tgcaaagcgg ggteeegtee 180
cgttttagcg gctctggatc cggcactgat tttaccctga ccattagcag cctgcaacct 240
gaagactttg cggtttatta ttgccagcag tatggtaata atcctactac ctttggccag 300
ggtacgaaag ttgaaattaa acgtacg 327
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Asp Arg Val Thr Ile Thr Cys Arg Ala Ser Gln Asn Ile Ser Asn Tyr 20 25 30
Leu Asn Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu Ile 35 40 45
Tyr Gly Thr Ser Ser Leu Gln Ser Gly Val Pro Ser Arg Phe Ser Gly 50 55 60
Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Gln Pro 65 70 75 80
Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Asn Asn Pro Thr 85 90 95
Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr 100 105
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cagtctcctg ggcgtggcct cgagtggctg ggccatatct attatcgtag caagtggtat 180
aactettatg eggtgagegt gaaaageegg attaceatea acceggatae ttegaaaaac 240

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cgttggggtg gtattcatga tggtgatatt tattttgatt attggggcca aggcaccctg	360
gtgacggtta gctcagc	377
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Thr Leu Ser Leu Thr Cys Ala Ile Ser Gly Asp Ser Val Ser Ser Asn 20 25 30	
Gly Ala Ala Trp Gly Trp Ile Arg Gln Ser Pro Gly Arg Gly Leu Glu 35 40 45	
Trp Leu Gly His Ile Tyr Tyr Arg Ser Lys Trp Tyr Asn Ser Tyr Ala	
Val Ser Val Lys Ser Arg Ile Thr Ile Asn Pro Asp Thr Ser Lys Asn 65 70 75 80	
Gln Phe Ser Leu Gln Leu Asn Ser Val Thr Pro Glu Asp Thr Ala Val	
85 90 95  Tyr Tyr Cys Ala Arg Trp Gly Gly Ile His Asp Gly Asp Ile Tyr Phe	
100 105 110	
Asp Tyr Trp Gly Gln Gly Thr Leu Val Thr Val Ser Ser 115 120 125	
<210> SEQ ID NO 113 <211> LENGTH: 327 <212> TYPE: DNA <213> ORGANISM: Homo sapiens	
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ggtaaagcac cgaaactatt aatttatgat gtttctaatt tgcaaagcgg ggtcccgtcc	180
cgttttagcg gctctggatc cggcactgat tttaccctga ccattagcag cctgcaacct	240
gaagactttg cggtttatta ttgccagcag tattctggtt atcctcttac ctttggccag	300
ggtacgaaag ttgaaattaa acgtacg	327
<210> SEQ ID NO 114 <211> LENGTH: 109 <212> TYPE: PRT <213> ORGANISM: Homo sapiens	
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Asp Arg Val Thr Ile Thr Cys Arg Ala Ser Gln Ser Ile Thr Asn Tyr	
Leu Asn Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu Ile	
Tyr Asp Val Ser Asn Leu Gln Ser Gly Val Pro Ser Arg Phe Ser Gly	
, , = ================================	

50 55 Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Gln Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Ser Gly Tyr Pro Leu Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr 100 <210> SEQ ID NO 115 <211> LENGTH: 362 <212> TYPE: DNA <213 > ORGANISM: Homo sapiens <400> SEOUENCE: 115 caggtgcaat tgcaacagtc tggtccgggc ctggtgaaac cgagccaaac cctgagcctg acctgtgcga tttccggaga tagcgtgagc tcttcttctg ctgcttggtc ttggattcgc cagtetectg ggegtggeet egagtggetg ggeatgatet attategtag caagtggtat aaccattatg cggtgagcgt gaaaagccgg attaccatca acccggatac ttcgaaaaac cagtttagcc tgcaactgaa cagcgtgacc ccggaagata cggccgtgta ttattgcgcg cgtggtggtt ctggtgttat ggatgtttgg ggccaaggca ccctggtgac ggttagctca <210> SEQ ID NO 116 <211> LENGTH: 120 <212> TYPE: PRT <213 > ORGANISM: Homo sapiens <400> SEQUENCE: 116 Gln Val Gln Leu Gln Gln Ser Gly Pro Gly Leu Val Lys Pro Ser Gln Thr Leu Ser Leu Thr Cys Ala Ile Ser Gly Asp Ser Val Ser Ser Ser Ser Ala Ala Trp Ser Trp Ile Arg Gln Ser Pro Gly Arg Gly Leu Glu Trp Leu Gly Met Ile Tyr Tyr Arg Ser Lys Trp Tyr Asn His Tyr Ala Val Ser Val Lys Ser Arg Ile Thr Ile Asn Pro Asp Thr Ser Lys Asn Gln Phe Ser Leu Gln Leu Asn Ser Val Thr Pro Glu Asp Thr Ala Val Tyr Tyr Cys Ala Arg Gly Gly Ser Gly Val Met Asp Val Trp Gly Gln 105 Gly Thr Leu Val Thr Val Ser Ser <210> SEQ ID NO 117 <211> LENGTH: 327 <212> TYPE: DNA <213 > ORGANISM: Homo sapiens <400> SEQUENCE: 117 60 gatatccaga tgacccagag cccgtctagc ctgagcgcga gcgtgggtga tcgtgtgacc attacctqca qaqcqaqcca qtctattaat ccttatctqa attqqtacca qcaqaaacca

ggtaaagcac cgaaactatt aatttatgct gcttctaatt tgcaaagcgg ggtcccgtcc cqttttaqcq qctctqqatc cqqcactqat tttaccctqa ccattaqcaq cctqcaacct gaagactttg cggtttatta ttgccagcag cttgataatc gttctattac ctttggccag 300 ggtacgaaag ttgaaattaa acgtacg 327 <210> SEQ ID NO 118 <211> LENGTH: 109 <212> TYPE: PRT <213> ORGANISM: Homo sapiens <400> SEOUENCE: 118 Asp Ile Gln Met Thr Gln Ser Pro Ser Ser Leu Ser Ala Ser Val Gly 10 Asp Arg Val Thr Ile Thr Cys Arg Ala Ser Gln Ser Ile Asn Pro Tyr 25 Leu Asn Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu Ile Tyr Ala Ala Ser Asn Leu Gln Ser Gly Val Pro Ser Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Gln Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Leu Asp Asn Arg Ser Ile Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr <210> SEQ ID NO 119 <211> LENGTH: 371 <212> TYPE: DNA <213 > ORGANISM: Homo sapiens <400> SEOUENCE: 119 caggtgcaat tgcaacagtc tggtccgggc ctggtgaaac cgagccaaac cctgagcctg 60 acctgtgcga tttccggaga tagcgtgagc tctaattctg ctgcttgggg ttggattcgc 120 cagtetectg ggegtggeet egagtggetg ggegttatet attategtag caagtggtat 180 aacgattatg cggtgagcgt gaaaagccgg attaccatca acccggatac ttcgaaaaac 240 cagtttagcc tgcaactgaa cagcgtgacc ccggaagata cggccgtgta ttattgcgcg 300 cgtgctcgtg ctaagaagtc tggtggtttt gattattggg gccaaggcac cctggtgacg 360 gttagctcag c 371 <210> SEQ ID NO 120 <211> LENGTH: 123 <212> TYPE: PRT <213> ORGANISM: Homo sapiens <400> SEQUENCE: 120 Gln Val Gln Leu Gln Gln Ser Gly Pro Gly Leu Val Lys Pro Ser Gln Thr Leu Ser Leu Thr Cys Ala Ile Ser Gly Asp Ser Val Ser Ser Asn Ser Ala Ala Trp Gly Trp Ile Arg Gln Ser Pro Gly Arg Gly Leu Glu

40

Trp Leu Gly Val Ile Tyr Tyr Arg Ser Lys Trp Tyr Asn Asp Tyr Ala Val Ser Val Lys Ser Arg Ile Thr Ile Asn Pro Asp Thr Ser Lys Asn Gln Phe Ser Leu Gln Leu Asn Ser Val Thr Pro Glu Asp Thr Ala Val Tyr Tyr Cys Ala Arg Ala Arg Ala Lys Lys Ser Gly Gly Phe Asp Tyr 105 Trp Gly Gln Gly Thr Leu Val Thr Val Ser Ser <210> SEQ ID NO 121 <211> LENGTH: 330 <212> TYPE: DNA <213 > ORGANISM: Homo sapiens <400> SEQUENCE: 121 gatatcgaac tgacccagcc gccttcagtg agcgttgcac caggtcagac cgcgcgtatc tcgtgtagcg gcgattctct tggttctaag tttgctcatt ggtaccagca gaaacccggg caggegecag ttettgtgat ttatgatgat tetaategte ceteaggeat eeeggaacge tttageggat ccaacagegg caacacegeg accetgacea ttageggeac tcaggeggaa gacgaagcgg attattattg ctctacttat acttctcgtt ctcattctta tgtgtttggc ggcggcacga agttaaccgt tcttggccag <210> SEQ ID NO 122 <211> LENGTH: 110 <212> TYPE: PRT <213 > ORGANISM: Homo sapiens <400> SEQUENCE: 122 Asp Ile Glu Leu Thr Gln Pro Pro Ser Val Ser Val Ala Pro Gly Gln Thr Ala Arg Ile Ser Cys Ser Gly Asp Ser Leu Gly Ser Lys Phe Ala His Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Val Leu Val Ile Tyr Asp Asp Ser Asn Arg Pro Ser Gly Ile Pro Glu Arg Phe Ser Gly Ser Asn Ser Gly Asn Thr Ala Thr Leu Thr Ile Ser Gly Thr Gln Ala Glu Asp Glu Ala Asp Tyr Tyr Cys Ser Thr Tyr Thr Ser Arg Ser His Ser Tyr Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu Gly Gln <210> SEQ ID NO 123 <211> LENGTH: 350 <212> TYPE: DNA <213 > ORGANISM: Homo sapiens <400> SEQUENCE: 123 caggtgcaat tggtggaaag cggcggcggc ctggtgcaac cgggcggcag cctgcgtctg agetgegegg ceteeggatt tacettttet tettatgett ettgggtgeg ceaageeect

gggaagggtc tcgagtgggt gagcggtatc tctggtgatg gtagcaatac ccattatgcg 180 qataqcqtqa aaqqccqttt taccatttca cqtqataatt cqaaaaacac cctqtatctq caaatgaaca gcctgcgtgc ggaagatacg gccgtgtatt attgcgcgcg ttatgataat 300 ttttattttg atgtttgggg ccaaggcacc ctggtgacgg ttagctcagc 350 <210> SEQ ID NO 124 <211> LENGTH: 116 <212> TYPE: PRT <213> ORGANISM: Homo sapiens <400> SEOUENCE: 124 Gln Val Gln Leu Val Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly 10 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Ser Tyr Ala Ser Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val Ser Gly Ile Ser Gly Asp Gly Ser Asn Thr His Tyr Ala Asp Ser Val Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys Ala Arg Tyr Asp Asn Phe Tyr Phe Asp Val Trp Gly Gln Gly Thr Leu Val 105 Thr Val Ser Ser 115 <210> SEQ ID NO 125 <211> LENGTH: 327 <212> TYPE: DNA <213 > ORGANISM: Homo sapiens <400> SEQUENCE: 125 gatategaac tgacceagec geetteagtg agegttgeac eaggteagac egegegtate 60 tcgtgtagcg gcgataatat tggttcttat tatgcttatt ggtaccagca gaaacccggg 120 caggegeeag ttettgtgat ttatgatgat tetaategte ceteaggeat eeeggaacge 180 tttageggat ccaacagegg caacacegeg accetgacca ttageggeac tcaggeggaa 240 gacgaagegg attattattg ceagtettat gattetaetg gtettettgt gtttggegge 300 ggcacgaagt taaccgttct tggccag 327 <210> SEQ ID NO 126 <211> LENGTH: 109 <212> TYPE: PRT <213 > ORGANISM: Homo sapiens <400> SEQUENCE: 126 Asp Ile Glu Leu Thr Gln Pro Pro Ser Val Ser Val Ala Pro Gly Gln Thr Ala Arg Ile Ser Cys Ser Gly Asp Asn Ile Gly Ser Tyr Tyr Ala Tyr Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Val Leu Val Ile Tyr

35 40 45	
Asp Asp Ser Asn Arg Pro Ser Gly Ile Pro Glu Arg Phe Ser Gly Ser 50 55 60	
Asn Ser Gly Asn Thr Ala Thr Leu Thr Ile Ser Gly Thr Gln Ala Glu 65 70 75 80	
Asp Glu Ala Asp Tyr Tyr Cys Gln Ser Tyr Asp Ser Thr Gly Leu Leu 85 90 95	
Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu Gly Gln 100 105	
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agctgcgcgg cctccggatt taccttttct aattatgcta tgacttgggt gcgccaagcc	120
cctgggaagg gtctcgagtg ggtgagcgtt atctcttctg ttggtagcaa tacctattat	180
geggatageg tgaaaggeeg ttttaceatt teaegtgata attegaaaaa caeeetgtat	240
ctgcaaatga acagcctgcg tgcggaagat acggccgtgt attattgcgc gcgtcctact	300
aaggotggto gtacttggtg gtggggtoot tatatggatg tttgggggooa aggoaccotg	360
gtgacggtta gctcagc	377
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Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Asn Tyr 20 25 30	
Ala Met Thr Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val 35 40 45	
Ser Val Ile Ser Ser Val Gly Ser Asn Thr Tyr Tyr Ala Asp Ser Val 50 55 60	
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr 65 70 75 80	
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys 85 90 95	
Ala Arg Pro Thr Lys Ala Gly Arg Thr Trp Trp Trp Gly Pro Tyr Met 100 105 110	
Asp Val Trp Gly Gln Gly Thr Leu Val Thr Val Ser Ser 115 120 125	
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caggogocag ttottgtgat ttatgatgat totaatogto cotcaggoat cooggaacgo	180								
tttageggat ccaacagegg caacacegeg accetgacea ttageggeac teaggeggaa	240								
gacgaagcgg attattattg cgagggttct aatgtgtttg gcggcggcac gaagttaacc	300								
gttettggee ag	312								
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Thr Ala Arg Ile Ser Cys Ser Gly Asp Asn Ile Gly Ser Tyr Phe Ala 20 25 30									
Ser Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Val Leu Val Ile Tyr 35 40 45									
Asp Asp Ser Asn Arg Pro Ser Gly Ile Pro Glu Arg Phe Ser Gly Ser 50 60									
Asn Ser Gly Asn Thr Ala Thr Leu Thr Ile Ser Gly Thr Gln Ala Glu 65 70 75 80									
Asp Glu Ala Asp Tyr Tyr Cys Glu Gly Ser Asn Val Phe Gly Gly 85 90 95									
Thr Lys Leu Thr Val Leu Gly Gln									
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cctgggaagg gtctcgagtg gatgggcatt atccagccgt ctgatagcga taccaattat	180								
teteegaget tteagggeea ggtgaecatt agegeggata aaageattag caeegegtat	240								
cttcaatgga gcagcctgaa agcgagcgat acggccatgt attattgcgc gcgttttatg	300								
tggtggggta agtatgattc tggttttgat gtttggggcc aaggcaccct ggtgacggtt	360								
agctcagc	368								
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Ser Leu Lys Ile Ser Cys Lys Gly Ser Gly Tyr Ser Phe Thr Asp Tyr 20 25 30

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Trp Ile Gly Trp Val Arg Gln Met Pro Gly Lys Gly Leu Glu Trp Met
                   40
Gly Ile Ile Gln Pro Ser Asp Ser Asp Thr Asn Tyr Ser Pro Ser Phe
Gln Gly Gln Val Thr Ile Ser Ala Asp Lys Ser Ile Ser Thr Ala Tyr
Leu Gln Trp Ser Ser Leu Lys Ala Ser Asp Thr Ala Met Tyr Tyr Cys
                                    90
Ala Arg Phe Met Trp Trp Gly Lys Tyr Asp Ser Gly Phe Asp Val Trp
Gly Gln Gly Thr Leu Val Thr Val Ser Ser
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caqqcqccaq ttcttqtqat ttatqqtqat aataatcqtc cctcaqqcat cccqqaacqc
tttaqcqqat ccaacaqcqq caacaccqcq accctqacca ttaqcqqcac tcaqqcqqaa
gacgaagcgg attattattg ccagtcttgg acttctcgtc ctatggttgt gtttggcggc
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                                                                     327
ggcacgaagt taaccgttct tggccag
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Thr Ala Arg Ile Ser Cys Ser Gly Asp Asn Leu Pro Ser Lys Ser Val
Tyr Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Val Leu Val Ile Tyr
                           40
Gly Asp Asn Asn Arg Pro Ser Gly Ile Pro Glu Arg Phe Ser Gly Ser
Asn Ser Gly Asn Thr Ala Thr Leu Thr Ile Ser Gly Thr Gln Ala Glu
                   70
Asp Glu Ala Asp Tyr Tyr Cys Gln Ser Trp Thr Ser Arg Pro Met Val
Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu Gly Gln
           100
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<213 > ORGANISM: Homo sapiens
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<221> NAME/KEY: misc_feature
<222> LOCATION: (156) .. (156)
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agetgegegg ceteeggatt tacettttet tettatteta tgeattgggt gegeeaagee
                                                                   120
cctgggaagg gtctcgagtg ggtgagcggt atctcntatt cttctagctt tacctattat
                                                                   180
gcggatagcg tgaaaggccg ttttaccatt tcacgtgata attcgaaaaa caccctgtat
                                                                   240
ctgcaaatga acagcctgcg tgcggaagat acggccgtgt attattgcgc gcgtgctctt
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ggtggtggtg ttgattattg gggccaaggc accctggtga cggttagctc agc
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Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Ser Tyr
Ser Met His Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
Ser Gly Ile Ser Tyr Ser Ser Ser Phe Thr Tyr Tyr Ala Asp Ser Val
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys
Val Thr Val Ser Ser
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<212> TYPE: DNA
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attacctgca gagcgagcca gggtatttct tcttatctgc attggtacca gcagaaacca
                                                                   120
ggtaaagcac cgaaactatt aatttatggt gcttctactt tgcaaagcgg ggtcccgtcc
                                                                   180
cgttttagcg gctctggatc cggcactgat tttaccctga ccattagcag cctgcaacct
                                                                   240
                                                                   300
gaagactttg cgacttatta ttgccagcag cagaatggtt atccttttac ctttggccag
                                                                   327
ggtacgaaag ttgaaattaa acgtacg
<210> SEQ ID NO 138
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<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 138
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Asp Arg Val Thr Ile Thr Cys Arg Ala Ser Gln Gly Ile Ser Ser Tyr 20 25 30										
Leu His Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu Ile 35 40 45										
Tyr Gly Ala Ser Thr Leu Gln Ser Gly Val Pro Ser Arg Phe Ser Gly 50 55 60										
Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Gln Pro 75 80										
Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Gln Asn Gly Tyr Pro Phe 85 90 95										
Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr 100 105										
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cagteteetg ggegtggeet egagtggetg ggeettatet attategtag caagtggtat 180										
aacgcttatg cggtgagcgt gaaaagccgg attaccatca acccggatac ttcgaaaaac 240										
cagtttagcc tgcaactgaa cagcgtgacc ccggaagata cggccgtgta ttattgcgcg 300										
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acggttagct cagc 374										
acggttagct cagc 374  <210> SEQ ID NO 140 <211> LENGTH: 124 <212> TYPE: PRT										
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acggttagct cagc 374  <210> SEQ ID NO 140 <211> LENGTH: 124 <212> TYPE: PRT <213> ORGANISM: Homo sapiens <400> SEQUENCE: 140  Gln Val Gln Leu Gln Gln Ser Gly Pro Gly Leu Val Lys Pro Ser Gln										
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                                                                     120
ggtaaagcac cgaaactatt aatttatgat gcttcttctt tgcaaagcgg ggtcccgtcc
cgttttagcg gctctggatc cggcactgat tttaccctga ccattagcag cctgcaacct
                                                                      240
gaagactttg cggtttatta ttgccagcag tattatgatt atcctcttac ctttggccag
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ggtacgaaag ttgaaattaa acgtacg
                                                                      327
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<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
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Asp Arg Val Thr Ile Thr Cys Arg Ala Ser Gln Asn Ile His Ser His
Leu Asn Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu Ile
Tyr Asp Ala Ser Ser Leu Gln Ser Gly Val Pro Ser Arg Phe Ser Gly
Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Gln Pro
Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Tyr Asp Tyr Pro Leu
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Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr
           100
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                                                                      120
cctgggaagg gtctcgagtg ggtgagctct atctcttctt cttctagcaa tacctattat
ggggatagcg tgaaaggccg ttttaccatt tcacgtgata attcgaaaaa caccctgtat
ctgcaaatga acagcctgcg tgcggaagat acggccgtgt attattgcgc gcgtatgcat
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<210> SEQ ID NO 144
<211> LENGTH: 117
<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 144
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<210> SEQ ID NO 147 <211> LENGTH: 350

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Ser Met Ser Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Va	1
35 40 45	_
Ser Ser Ile Ser Ser Ser Ser Ser Asn Thr Tyr Tyr Gly Asp Ser Va 50 60	1
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Ty 65 70 75 80	
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cy 85 90 95	ន
Ala Arg Met His Tyr Lys Gly Met Asp Ile Trp Gly Gln Gly Thr Le	u
Val Thr Val Ser Ser 115	
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caggogocag ttottgtgat ttatggtgat totaagogto cotcaggoat cooggaa	cgc 180
tttageggat ecaacagegg caacacegeg accetgacea ttageggeac teaggeg	gaa 240
gacgaagegg attattattg etettetget gettttggtt etaetgtgtt tggegge	ggc 300
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Asp Ile Glu Leu Thr Gln Pro Pro Ser Val Ser Val Ala Pro Gly Gl 1 5 10 15	n
Thr Ala Arg Ile Ser Cys Ser Gly Asp Lys Leu Gly Lys Tyr Tyr Al 20 25 30	a
Tyr Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Val Leu Val Ile Ty 35 40 45	r
Gly Asp Ser Lys Arg Pro Ser Gly Ile Pro Glu Arg Phe Ser Gly Se 50 55 60	r
Asn Ser Gly Asn Thr Ala Thr Leu Thr Ile Ser Gly Thr Gln Ala Gl 65 70 75 80	
Asp Glu Ala Asp Tyr Tyr Cys Ser Ser Ala Ala Phe Gly Ser Thr Va 85 90 95	1
Phe Gly Gly Gly Thr Lys Leu Thr Val Leu Gly Gln	

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<212> TYPE: DNA
<213 > ORGANISM: Homo sapiens
<400> SEOUENCE: 147
caggtgcaat tggtggaaag cggcggcggc ctggtgcaac cgggcggcag cctgcgtctg
                                                                      60
agetgegegg ceteeggatt tacetttaat tettattata tgtettgggt gegeeaagee
                                                                     120
cctgggaagg gtctcgagtg ggtgagcaat atctcttctt ctggtagcaa taccaattat
                                                                     180
gcggatagcg tgaaaggccg ttttaccatt tcacgtgata attcgaaaaa caccctgtat
ctgcaaatga acagcctgcg tgcggaagat acggccgtgt attattgcgc gcgtgttcat
                                                                     300
tatggttttg atttttgggg ccaaggcacc ctggtgacgg ttagctcagc
                                                                     350
<210> SEQ ID NO 148
<211> LENGTH: 116
<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 148
Gln Val Gln Leu Val Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Asn Ser Tyr
Tyr Met Ser Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
Ser Asn Ile Ser Ser Ser Gly Ser Asn Thr Asn Tyr Ala Asp Ser Val
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys
Ala Arg Val His Tyr Gly Phe Asp Phe Trp Gly Gln Gly Thr Leu Val
                               105
Thr Val Ser Ser
       115
<210> SEO ID NO 149
<211> LENGTH: 327
<212> TYPE: DNA
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 149
gatategaac tgacceagee geetteagtg agegttgeac caggteagac egegegtate
                                                                      60
tcgtgtagcg gcgatgctct tggttctaag tttgctcatt ggtaccagca gaaacccggg
caggegecag ttettgtgat ttatgatgat tetgagegte ceteaggeat eeeggaacge
tttageggat ccaacagegg caacacegeg accetgacea ttageggeac tcaggeggaa
gacgaagcgg attattattg ccaggettat gattetggte ttetttatgt gtttggegge
ggcacgaagt taaccgttct tggccag
                                                                      327
<210> SEQ ID NO 150
<211> LENGTH: 109
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 150
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Asp Ile Glu Leu Thr Gln Pro Pro Ser Val Ser Val Ala Pro Gly Gln Thr Ala Arg Ile Ser Cys Ser Gly Asp Ala Leu Gly Ser Lys Phe Ala His Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Val Leu Val Ile Tyr Asp Asp Ser Glu Arg Pro Ser Gly Ile Pro Glu Arg Phe Ser Gly Ser Asn Ser Gly Asn Thr Ala Thr Leu Thr Ile Ser Gly Thr Gln Ala Glu Asp Glu Ala Asp Tyr Tyr Cys Gln Ala Tyr Asp Ser Gly Leu Leu Tyr Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu Gly Gln <210> SEQ ID NO 151 <211> LENGTH: 359 <212> TYPE: DNA <213 > ORGANISM: Homo sapiens <400> SEQUENCE: 151 caggtgcaat tggtggaaag cggcggcggc ctggtgcaac cgggcggcag cctgcgtctg agctgcgcgg cctccggatt tacctttcgt aattatgcta tgaattgggt gcgccaagcc cctgggaagg gtctcgagtg ggtgagcgtt atctctggtt cttctagcta tacctattat geggatageg tgaaaggeeg ttttaccatt teaegtgata attegaaaaa caccetgtat ctgcaaatga acagcctgcg tgcggaagat acggccgtgt attattgcgc gcgtgctgat 300 cttccttata tggtttttga ttattggggc caaggcaccc tggtgacggt tagctcagc <210> SEO ID NO 152 <211> LENGTH: 119 <212> TYPE: PRT <213 > ORGANISM: Homo sapiens <400> SEQUENCE: 152 Gln Val Gln Leu Val Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Arg Asn Tyr Ala Met Asn Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val Ser Val Ile Ser Gly Ser Ser Ser Tyr Thr Tyr Tyr Ala Asp Ser Val 55 Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys Ala Arg Ala Asp Leu Pro Tyr Met Val Phe Asp Tyr Trp Gly Gln Gly Thr Leu Val Thr Val Ser Ser

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<211> LENGTH: 324
<212> TYPE: DNA
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 153
gatategaac tgacceagee geetteagtg agegttgeac caggteagae egegegtate
                                                                       60
tcgtgtagcg gcgatgctct tggtaagtat tatgcttctt ggtaccagca gaaacccggg
                                                                     120
caggegecag ttettgtgat ttatggtgat aataagegte ceteaggeat eeeggaacge
                                                                     180
tttageggat ccaacagegg caacacegeg accetgacca ttageggcae tcaggeggaa
                                                                     240
gacgaagcgg attattattg ccagtcttat actactcgtt ctcttgtgtt tggcggcggc
                                                                     300
acgaagttaa ccgttcttgg ccag
                                                                     324
<210> SEQ ID NO 154
<211> LENGTH: 108
<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 154
Asp Ile Glu Leu Thr Gln Pro Pro Ser Val Ser Val Ala Pro Gly Gln
Thr Ala Arg Ile Ser Cys Ser Gly Asp Ala Leu Gly Lys Tyr Tyr Ala
Ser Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Val Leu Val Ile Tyr
Gly Asp Asn Lys Arg Pro Ser Gly Ile Pro Glu Arg Phe Ser Gly Ser
Asn Ser Gly Asn Thr Ala Thr Leu Thr Ile Ser Gly Thr Gln Ala Glu
Asp Glu Ala Asp Tyr Tyr Cys Gln Ser Tyr Thr Thr Arg Ser Leu Val
Phe Gly Gly Gly Thr Lys Leu Thr Val Leu Gly Gln
<210> SEQ ID NO 155
<211> LENGTH: 353
<212> TYPE: DNA
<213 > ORGANISM: Homo sapiens
<400> SEOUENCE: 155
caggtgcaat tggtggaaag cggcggcggc ctggtgcaac cgggcggcag cctgcgtctg
                                                                       60
agetgegegg ceteeggatt tacettttet tettatggta tgtettgggt gegeeaagee
                                                                     120
cctgggaagg gtctcgagtg ggtgagcctt atctctggtg tttctagctc tacctattat
geggatageg tgaaaggeeg ttttaccatt teaegtgata attegaaaaa caccetgtat
ctgcaaatga acagcctgcg tgcggaagat acggccgtgt attattgcgc gcgttcttat
cttggttatt ttgatgtttg gggccaaggc accctggtga cggttagctc agc
                                                                      353
<210> SEQ ID NO 156
<211> LENGTH: 117
<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 156
Gln Val Gln Leu Val Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
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1 5 10 15	
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Ser Tyr 20 25 30	
Gly Met Ser Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val 35 40 45	
Ser Leu Ile Ser Gly Val Ser Ser Ser Thr Tyr Tyr Ala Asp Ser Val 50 55 60	
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr 65 70 75 80	
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys 85 90 95	
Ala Arg Ser Tyr Leu Gly Tyr Phe Asp Val Trp Gly Gln Gly Thr Leu 100 105 110	
Val Thr Val Ser Ser 115	
<210> SEQ ID NO 157 <211> LENGTH: 342 <212> TYPE: DNA <213> ORGANISM: Homo sapiens	
<400> SEQUENCE: 157	
gatategtga tgacceagag eccaetgage etgecagtga etcegggega geetgegage	60
attagctgca gaagcagcca aagcctggtt ttttctgatg gcaatactta tctgaattgg	120
taccttcaaa aaccaggtca aagcccgcag ctattaattt ataagggttc taatcgtgcc	180
agtggggtcc cggatcgttt tagcggctct ggatccggca ccgattttac cctgaaaatt	240
agccgtgtgg aagctgaaga cgtgggcgtg tattattgcc agcagtatga ttcttatcct	300
cttacctttg gccagggtac gaaagttgaa attaaacgta cg	342
<210> SEQ ID NO 158 <211> LENGTH: 114 <212> TYPE: PRT <213> ORGANISM: Homo sapiens	
<400> SEQUENCE: 158	
Asp Ile Val Met Thr Gln Ser Pro Leu Ser Leu Pro Val Thr Pro Gly 1 10 15	
Glu Pro Ala Ser Ile Ser Cys Arg Ser Ser Gln Ser Leu Val Phe Ser 20 25 30	
Asp Gly Asn Thr Tyr Leu Asn Trp Tyr Leu Gln Lys Pro Gly Gln Ser 35 40 45	
Pro Gln Leu Leu Ile Tyr Lys Gly Ser Asn Arg Ala Ser Gly Val Pro 50 60	
Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Lys Ile 65 70 75 80	
Ser Arg Val Glu Ala Glu Asp Val Gly Val Tyr Tyr Cys Gln Gln Tyr 85 90 95	
Asp Ser Tyr Pro Leu Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys	
Arg Thr	

<211> LENGTH: 371 <212> TYPE: DNA <213> ORGANISM: Homo sapiens	
<400> SEQUENCE: 159	
caggtgcaat tgcaacagtc tggtccgggc ctggtgaaac cgagccaaac cctgagcctg	60
acctgtgcga tttccggaga tagcgtgagc tctaattctg ctgcttggtc ttggattcgc	120
cagtctcctg ggcgtggcct cgagtggctg ggcattatct ataagcgtag caagtggtat	180
aacgattatg cggtgagcgt gaaaagccgg attaccatca acccggatac ttcgaaaaac	240
cagtttagcc tgcaactgaa cagcgtgacc ccggaagata cggccgtgta ttattgcgcg	300
cgttggcatt ctgataagca ttggggtttt gattattggg gccaaggcac cctggtgacg	360
gttagctcag c	371
<210> SEQ ID NO 160 <211> LENGTH: 123 <212> TYPE: PRT <213> ORGANISM: Homo sapiens	
<400> SEQUENCE: 160	
Gln Val Gln Leu Gln Gln Ser Gly Pro Gly Leu Val Lys Pro Ser Gln 1 5 15	
Thr Leu Ser Leu Thr Cys Ala Ile Ser Gly Asp Ser Val Ser Ser Asn 20 25 30	
Ser Ala Ala Trp Ser Trp Ile Arg Gln Ser Pro Gly Arg Gly Leu Glu 35 40 45	
Trp Leu Gly Ile Ile Tyr Lys Arg Ser Lys Trp Tyr Asn Asp Tyr Ala 50 55 60	
Val Ser Val Lys Ser Arg Ile Thr Ile Asn Pro Asp Thr Ser Lys Asn 65 70 75 80	
Gln Phe Ser Leu Gln Leu Asn Ser Val Thr Pro Glu Asp Thr Ala Val 85 90 95	
Tyr Tyr Cys Ala Arg Trp His Ser Asp Lys His Trp Gly Phe Asp Tyr 100 105 110	
Trp Gly Gln Gly Thr Leu Val Thr Val Ser Ser 115 120	
<210> SEQ ID NO 161 <211> LENGTH: 327 <212> TYPE: DNA <213> ORGANISM: Homo sapiens	
<400> SEQUENCE: 161	
gatatogaac tgacocagoo goottoagtg agogttgcac caggtoagac ogogogtato	60
tegtgtageg gegatgetet tggttetaag tatgtttett ggtaceagea gaaaceeggg	120
caggogocag ttottgtgat ttatggtgat aataagogto ootcaggoat oooggaaogo	180
tttagoggat ocaacagogg caacacogog accotgacoa ttagoggoac tcaggoggaa	240
gacgaagcgg attattattg ccagtcttat acttattctc ttaatcaggt gtttggcggc	300
ggcacgaagt taaccgttct tggccag	327
<210> SEQ ID NO 162 <211> LENGTH: 109 <212> TYPE: PRT	

<sup>&</sup>lt;212> TYPE: PRT

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<213 > ORGANISM: Homo sapiens
<400> SEOUENCE: 162
Asp Ile Glu Leu Thr Gln Pro Pro Ser Val Ser Val Ala Pro Gly Gln
                                   10
Thr Ala Arg Ile Ser Cys Ser Gly Asp Ala Leu Gly Ser Lys Tyr Val
Ser Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Val Leu Val Ile Tyr
Gly Asp Asn Lys Arg Pro Ser Gly Ile Pro Glu Arg Phe Ser Gly Ser
Asn Ser Gly Asn Thr Ala Thr Leu Thr Ile Ser Gly Thr Gln Ala Glu
Asp Glu Ala Asp Tyr Tyr Cys Gln Ser Tyr Thr Tyr Ser Leu Asn Gln
Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu Gly Gln
<210> SEQ ID NO 163
<211> LENGTH: 356
<212> TYPE: DNA
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 163
caqqtqcaat tqqtqqaaaq cqqcqqcqqc ctqqtqcaac cqqqcqqcaq cctqcqtctq
                                                                     120
agetgegegg ceteeggatt tacetttaat gattatgeta tgtettgggt gegeeaagee
cctqqqaaqq qtctcqaqtq qqtqaqcctt atcqaqtctq tttctaqctc tacctattat
                                                                     180
qcqqataqcq tqaaaqqccq ttttaccatt tcacqtqata attcqaaaaa caccctqtat
                                                                     240
ctgcaaatga acagcctgcg tgcggaagat acggccgtgt attattgcgc gcgtactatt
                                                                     300
ggtgttcttt gggatgatgt ttggggccaa ggcaccctgg tgacggttag ctcagc
                                                                     356
<210> SEO ID NO 164
<211> LENGTH: 118
<212> TYPE: PRT
<213> ORGANISM: Homo sapiens
<400> SEQUENCE: 164
Gln Val Gln Leu Val Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
                                   10
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Asn Asp Tyr
                               25
Ala Met Ser Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
                           40
Ser Leu Ile Glu Ser Val Ser Ser Ser Thr Tyr Tyr Ala Asp Ser Val
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys
Ala Arg Thr Ile Gly Val Leu Trp Asp Asp Val Trp Gly Gln Gly Thr
Leu Val Thr Val Ser Ser
       115
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<210> SEQ ID NO 165
<211> LENGTH: 324
<212> TYPE: DNA
<213> ORGANISM: Homo sapiens
<400> SEQUENCE: 165
gatategaac tgacceagee geetteagtg agegttgeac caggteagae egegegtate
                                                                      60
tcgtgtagcg gcgataagct tggttctaag tctgttcatt ggtaccagca gaaacccggg
                                                                     120
caggegeeag ttettgtgat ttategtgat aetgategte ceteaggeat eeeggaaege
                                                                     180
tttagcggat ccaacagcgg caacaccgcg accctgacca ttagcggcac tcaggcggaa
                                                                     240
gacgaagcgg attattattg ccagacttat gattatattc ttaatgtgtt tggcggcggc
                                                                     300
acgaagttaa ccgttcttgg ccag
                                                                     324
<210> SEQ ID NO 166
<211> LENGTH: 108
<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 166
Asp Ile Glu Leu Thr Gln Pro Pro Ser Val Ser Val Ala Pro Gly Gln
Thr Ala Arg Ile Ser Cys Ser Gly Asp Lys Leu Gly Ser Lys Ser Val
His Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Val Leu Val Ile Tyr
Arg Asp Thr Asp Arg Pro Ser Gly Ile Pro Glu Arg Phe Ser Gly Ser
Asn Ser Gly Asn Thr Ala Thr Leu Thr Ile Ser Gly Thr Gln Ala Glu
Asp Glu Ala Asp Tyr Tyr Cys Gln Thr Tyr Asp Tyr Ile Leu Asn Val
Phe Gly Gly Gly Thr Lys Leu Thr Val Leu Gly Gln
<210> SEQ ID NO 167
<211> LENGTH: 359
<212> TYPE: DNA
<213> ORGANISM: Homo sapiens
<400> SEQUENCE: 167
caggtgcaat tggtggaaag cggcggcggc ctggtgcaac cgggcggcag cctgcgtctg
agetgegegg ceteeggatt tacettttet aettatgeta tgeattgggt gegeeaagee
                                                                     120
cctgggaagg gtctcgagtg ggtgagcact atctctggtt atggtagctt tacctattat
geggatageg tgaaaggeeg ttttaccatt teaegtgata attegaaaaa caccetgtat
ctgcaaatga acagcctgcg tgcggaagat acggccgtgt attattgcgc gcgtaatggt
cgtaagtatg gtcagatgga taattggggc caaggcaccc tggtgacggt tagctcagc
<210> SEQ ID NO 168
<211> LENGTH: 119
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<sup>&</sup>lt;212> TYPE: PRT

<sup>&</sup>lt;213 > ORGANISM: Homo sapiens

< 400	)> SI	EQUE	ICE :	168												
Gln 1	Val	Gln	Leu	Val 5	Glu	Ser	Gly	Gly	Gly 10	Leu	Val	Gln	Pro	Gly 15	Gly	
Ser	Leu	Arg	Leu 20	Ser	Cys	Ala	Ala	Ser 25	Gly	Phe	Thr	Phe	Ser 30	Thr	Tyr	
Ala	Met	His 35	Trp	Val	Arg	Gln	Ala 40	Pro	Gly	Lys	Gly	Leu 45	Glu	Trp	Val	
Ser	Thr 50	Ile	Ser	Gly	Tyr	Gly 55	Ser	Phe	Thr	Tyr	Tyr 60	Ala	Asp	Ser	Val	
Lys 65	Gly	Arg	Phe	Thr	Ile 70	Ser	Arg	Asp	Asn	Ser 75	Lys	Asn	Thr	Leu	Tyr 80	
Leu	Gln	Met	Asn	Ser 85	Leu	Arg	Ala	Glu	Asp 90	Thr	Ala	Val	Tyr	Tyr 95	Cys	
Ala	Arg	Asn	Gly 100	Arg	Lys	Tyr	Gly	Gln 105	Met	Asp	Asn	Trp	Gly 110	Gln	Gly	
Thr	Leu	Val 115	Thr	Val	Ser	Ser										
<210> SEQ ID NO 169 <211> LENGTH: 333 <212> TYPE: DNA <213> ORGANISM: Homo sapiens																
< 400	)> SI	EQUE	ICE :	169												
gata	atcga	aac t	gaco	ccago	ec go	ctto	cagto	g ago	gttg	gcac	cago	gtcaç	gac o	gege	gtatc	60
tcgt	gtag	geg g	gcgat	tcta	at to	gtaa	agaaç	g tat	gtto	att	ggta	accaç	gca q	gaaac	ccggg	120
cag	geged	ag t	tctt	gtga	at tt	atg	gtgat	aat	aato	gtc	ccto	caggo	cat o	cccgc	gaacgc	180
ttta	agegg	gat o	caac	cagco	gg ca	acac	ccgcc	g acc	cctga	acca	ttag	gegge	cac t	cago	gcggaa	240
gac	gaago	gg a	attat	tatt	g ct	ctac	ctgct	gat	tctç	gtta	ttad	cttat	aa q	gaato	gtgttt	300
ggc	ggcgg	gca c	gaag	gttaa	ac co	gttct	tggo	caç	3							333
<210> SEQ ID NO 170 <211> LENGTH: 111 <212> TYPE: PRT <213> ORGANISM: Homo sapiens <400> SEQUENCE: 170																
Asp 1	Ile	Glu	Leu	Thr 5	Gln	Pro		Ser		Ser	Val	Ala	Pro	Gly 15	Gln	
Thr	Ala	Arg	Ile 20	Ser	Cys	Ser	Gly	Asp 25	Ser	Ile	Gly	Lys	Lys 30	Tyr	Val	
His	Trp	Tyr 35	Gln	Gln	Lys	Pro	Gly 40	Gln	Ala	Pro	Val	Leu 45	Val	Ile	Tyr	
Gly	Asp 50	Asn	Asn	Arg	Pro	Ser 55	Gly	Ile	Pro	Glu	Arg 60	Phe	Ser	Gly	Ser	
Asn 65	Ser	Gly	Asn	Thr	Ala 70	Thr	Leu	Thr	Ile	Ser 75	Gly	Thr	Gln	Ala	Glu 80	
Asp	Glu	Ala	Asp	Tyr 85	Tyr	Cys	Ser	Thr	Ala 90	Asp	Ser	Val	Ile	Thr 95	Tyr	
Lys	Asn	Val	Phe 100	Gly	Gly	Gly	Thr	Lys 105	Leu	Thr	Val	Leu	Gly 110	Gln		

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<210> SEQ ID NO 171
<211> LENGTH: 362
<212> TYPE: DNA
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 171
caggtgcaat tggtggaaag cggcggcggc ctggtgcaac cgggcggcag cctgcgtctg
                                                                      60
agetgegegg ceteeggatt tacettttet gateatgeta tgeattgggt gegeeaagee
cctgggaagg gtctcgagtg ggtgagcgtt atcgagtatt ctggtagcaa gaccaattat
geggatageg tgaaaggeeg ttttaccatt teaegtgata attegaaaaa caccetgtat
                                                                     240
ctgcaaatga acagcctgcg tgcggaagat acggccgtgt attattgcgc gcgtggtgat
                                                                     300
tattatcctt atcttgtttt tgctatttgg ggccaaggca ccctggtgac ggttagctca
                                                                       362
<210> SEQ ID NO 172
<211> LENGTH: 120
<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 172
Gln Val Gln Leu Val Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly 1 \phantom{\bigg|} 5 \phantom{\bigg|} 10 \phantom{\bigg|} 15
Ala Met His Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
Ser Val Ile Glu Tyr Ser Gly Ser Lys Thr Asn Tyr Ala Asp Ser Val
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr
                   70
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys
Ala Arg Gly Asp Tyr Tyr Pro Tyr Leu Val Phe Ala Ile Trp Gly Gln
           100
                               105
Gly Thr Leu Val Thr Val Ser Ser
      115
<210> SEO ID NO 173
<211> LENGTH: 11 <212> TYPE: PRT
<213> ORGANISM: Homo sapiens
<400> SEQUENCE: 173
Ser Gly Asp Asn Ile Arg Thr Tyr Tyr Val His
    5
<210> SEQ ID NO 174
<211> LENGTH: 11
<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 174
Ser Gly Asp Asn Ile Pro Glu Lys Tyr Val His
```

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<210> SEQ ID NO 175
<211> LENGTH: 11
<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 175
Ser Gly Asp Lys Ile Gly Ser Lys Tyr Val Tyr
              5
<210> SEQ ID NO 176
<211> LENGTH: 11
<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 176
Ser Gly Asp Asn Leu Arg Asn Tyr Tyr Ala His
        5
<210> SEQ ID NO 177
<211> LENGTH: 11
<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 177
Ser Gly Asp Lys Leu Gly Lys Lys Tyr Val His
<210> SEQ ID NO 178
<211> LENGTH: 11
<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 178
Ser Gly Asp Asn Leu Gly Asn Lys Tyr Ala His
              5
<210> SEQ ID NO 179
<211> LENGTH: 12
<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 179
Arg Ala Ser Gln Asn Ile Gly Ser Asn Tyr Leu Ala
1 5
<210> SEQ ID NO 180
<211> LENGTH: 11
<212> TYPE: PRT
<213> ORGANISM: Homo sapiens
<400> SEQUENCE: 180
Ser Gly Asp Ala Leu Gly Thr Tyr Tyr Ala Tyr
1 5
<210> SEQ ID NO 181
<211> LENGTH: 11
<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 181
Ser Gly Asp Asn Leu Arg Gly Tyr Tyr Ala Ser
               5
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<210> SEO ID NO 182
<211> LENGTH: 11
<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 182
Arg Ala Ser Gln Ser Ile Arg Ser Tyr Leu Ala
<210> SEQ ID NO 183
<211> LENGTH: 11
<212> TYPE: PRT
<213> ORGANISM: Homo sapiens
<400> SEQUENCE: 183
Ser Gly Asp Ser Ile Gly Ser Tyr Tyr Ala His
<210> SEQ ID NO 184
<211> LENGTH: 11
<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 184
Arg Ala Ser Gln Asp Ile Arg Asn Asn Leu Ala
1 5
<210> SEQ ID NO 185
<211> LENGTH: 16
<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 185
Arg Ser Ser Gln Ser Leu Leu His Ser Asn Gly Tyr Thr Tyr Leu Ser
               5
                                    10
<210> SEQ ID NO 186
<211> LENGTH: 14
<212> TYPE: PRT
<213> ORGANISM: Homo sapiens
<400> SEQUENCE: 186
Thr Gly Thr Ser Ser Asp Ile Gly Gly Tyr Asn Tyr Val Ser 1 \phantom{\bigg|}
<210> SEQ ID NO 187
<211> LENGTH: 11
<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 187
Arg Ala Ser Gln Pro Ile Tyr Asn Ser Leu Ser
     5
<210> SEQ ID NO 188
<211> LENGTH: 12
<212> TYPE: PRT
<213 > ORGANISM: Homo sapiens
<400> SEQUENCE: 188
Arg Ala Ser Gln Ser Val Ser Ser Gln Tyr Leu Ala
```

```
<210> SEQ ID NO 189
<211> LENGTH: 11
<212> TYPE: PRT
<213> ORGANISM: Homo sapiens
<400> SEQUENCE: 189
Ser Gly Asp Asn Ile Arg Lys Phe Tyr Val His
1 5
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<211> LENGTH: 11
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Ser Tyr Ser Met His
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Asp His Ala Met His
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Thr Ile Ser Tyr Asp Gly Ser Asn Thr Tyr Tyr Ala Asp Ser Val Lys
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Gly
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Gly
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Lys Ser
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Lys Ser
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Lys Ser
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Lys Ser
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Lys Ser
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Lys Ser
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Lys Ser
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Lys Ser
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Lys Ser
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Lys Ser
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Lys Ser
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Lys Ser
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Gly Val His Ser Tyr Phe Asp Tyr
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Tyr Tyr Cys Lys Cys Val Asp Leu
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Lys Gly Gly Glu His Gly Phe Phe Pro Ser Asp Ile
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Tyr Gly Gly Met Asp Tyr
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Ala Arg Ala Lys Lys Ser Gly Gly Phe Asp Tyr
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Tyr Asp Asn Phe Tyr Phe Asp Val
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# We claim:

- 1. An isolated monoclonal antibody that binds to human tissue factor pathway inhibitor, wherein the antibody comprises a CDR3 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 388-430.
- 2. The isolated antibody of claim 1, wherein the antibody further comprises (a) a CDR1 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 302-344, (b) a CDR2 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 345-387, or (c) both a CDR1 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 302-344 and a CDR2 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 345-387.
- 3. An isolated monoclonal antibody that binds to human tissue factor pathway inhibitor, wherein the antibody comprises a CDR3 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 259-301.
- **4.** The isolated antibody of claim **3**, wherein the antibody further comprises (a) a CDR1 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 173-215, (b) a CDR2 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 216-258, or (c) both a CDR1 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 173-215 and a CDR2 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 216-258.
- 5. The antibody of claim 1, wherein the antibody further comprises a CDR3 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 259-301.
- 6. The antibody of claim 5, wherein the antibody further comprises (a) a CDR1 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 302-344, (b) a CDR2 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 345-387, (c) a CDR1 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 173-215, and (d) a CDR2 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 216-258.
- 7. The antibody of claim 1, wherein the antibody comprises heavy and light chain variable regions comprising:
  - (a) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 173, 216 and 259 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 302, 345 and 388;
  - (b) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 174, 217 and 260

- and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 303, 346 and 389;
- (c) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 175, 218 and 261 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 304, 347 and 390;
- (d) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 176, 219 and 262 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 305, 348 and 391;
- (e) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 177, 220 and 263 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 306, 349 and 392:
- (f) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 178, 221 and 264 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 307, 350 and 393;
- (g) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 179, 222 and 265 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 308, 351 and 304.
- (h) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 180, 223 and 266 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 309, 352 and 305.
- (i) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 181, 224 and 267 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 310, 353 and 396;
- (j) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 182, 225 and 268 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 311, 354 and 397;
- (k) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 183, 226 and 269 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 312, 355 and 398;

- (1) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 184, 227 and 270 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 313, 356 and 399;
- (m) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 185, 228 and 271 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 314, 357 and 400:
- (n) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 186, 229 and 272 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 315, 358 and 401;
- (o) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 187, 230 and 273 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 316, 359 and 402:
- (p) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 188, 231 and 274 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 317, 360 and 403;
- (q) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 189, 232 and 275 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 318, 361 and 404.
- (r) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 190, 233 and 276 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 319, 362 and 405;
- (s) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 191, 234 and 277 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 320, 363 and 406;
- (t) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 192, 235 and 278 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 321, 364 and 407:
- (u) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 193, 236 and 279 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 322, 365 and 408:
- (v) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 194, 237 and 280 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 323, 366 and 409;
- (w) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 195, 238 and 281 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 324, 367 and 410;
- (x) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 196, 239 and 282

- and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 325, 368 and 411:
- (y) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 197, 240 and 283 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 326, 369 and 412:
- (z) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 198, 241 and 284 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 327, 370 and 413:
- (aa) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 199, 242 and 285 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 328, 371 and 414:
- (bb) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 200, 243 and 286 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 329, 372 and 415.
- (cc) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 201, 244 and 287 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 330, 373 and;
- (dd) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 202, 245 and 288 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 331, 374 and 417;
- (ee) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 203, 246 and 289 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 332, 375 and 418;
- (ff) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 204, 247 and 290 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 333, 376 and 419:
- (gg) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 205, 248 and 291 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 334, 377 and 420;
- (hh) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 206, 249 and 292 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 335, 378 and 421.
- (ii) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 207, 250 and 293 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 336, 379 and 422;
- (jj) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 208, 251 and 294 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 337, 380 and 423:
- (kk) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 209, 252 and 295

- and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 338, 381 and 424:
- (II) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 210, 253 and 296 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 339, 382 and 425:
- (mm) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 211, 254 and 297 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 340, 383 and 426;
- (nn) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 212, 255 and 298 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 341, 384 and 427:
- (oo) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 213, 256 and 299 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 342, 385 and 428.
- (pp) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 214, 257 and 300 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 343, 386 and 429;
- (qq) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 215, 258 and 301 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 344, 387 and 430; or
- (rr) a light chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 194, 237 and 280 and a heavy chain variable region comprising an amino acid sequence comprising SEQ ID NOs: 335, 378 and 421.
- 8. The monoclonal antibody of claim 1, comprising:
- (a) a light chain variable region having the polypeptide sequence of SEQ ID NO: 2 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 4.
- (b) a light chain variable region having the polypeptide sequence of SEQ ID NO: 6 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 8;
- (c) a light chain variable region having the polypeptide sequence of SEQ ID NO: 10 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 12:
- (d) a light chain variable region having the polypeptide sequence of SEQ ID NO: 14 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 16;
- (e) a light chain variable region having the polypeptide sequence of SEQ ID NO: 18 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 20;
- (f) a light chain variable region having the polypeptide sequence of SEQ ID NO: 22 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 24:

- (g) a light chain variable region having the polypeptide sequence of SEQ ID NO: 26 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 28:
- (h) a light chain variable region having the polypeptide sequence of SEQ ID NO: 30 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 32.
- (i) a light chain variable region having the polypeptide sequence of SEQ ID NO: 34 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 26.
- (j) a light chain variable region having the polypeptide sequence of SEQ ID NO: 38 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 40.
- (k) a light chain variable region having the polypeptide sequence of SEQ ID NO: 42 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 44;
- a light chain variable region having the polypeptide sequence of SEQ ID NO: 46 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 48;
- (m) a light chain variable region having the polypeptide sequence of SEQ ID NO: 50 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 52:
- (n) a light chain variable region having the polypeptide sequence of SEQ ID NO: 54 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 56.
- (o) a light chain variable region having the polypeptide sequence of SEQ ID NO: 58 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO:
- (p) a light chain variable region having the polypeptide sequence of SEQ ID NO: 62 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 64;
- (q) a light chain variable region having the polypeptide sequence of SEQ ID NO: 66 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 68;
- (r) a light chain variable region having the polypeptide sequence of SEQ ID NO: 70 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 72:
- (s) a light chain variable region having the polypeptide sequence of SEQ ID NO: 74 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 76:
- (t) a light chain variable region having the polypeptide sequence of SEQ ID NO: 78 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 80.
- (u) a light chain variable region having the polypeptide sequence of SEQ ID NO: 82 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 84.
- (v) a light chain variable region having the polypeptide sequence of SEQ ID NO: 86 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 88.

- (w) a light chain variable region having the polypeptide sequence of SEQ ID NO: 90 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 92:
- (x) a light chain variable region having the polypeptide sequence of SEQ ID NO: 94 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 96.
- (y) a light chain variable region having the polypeptide sequence of SEQ ID NO: 98 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 100.
- (z) a light chain variable region having the polypeptide sequence of SEQ ID NO: 102 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 104:
- (aa) a light chain variable region having the polypeptide sequence of SEQ ID NO: 106 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 108;
- (bb) a light chain variable region having the polypeptide sequence of SEQ ID NO: 110 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 112:
- (cc) a light chain variable region having the polypeptide sequence of SEQ ID NO: 114 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 116:
- (dd) a light chain variable region having the polypeptide sequence of SEQ ID NO: 118 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 120;
- (ee) a light chain variable region having the polypeptide sequence of SEQ ID NO: 122 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 124:
- (ff) a light chain variable region having the polypeptide sequence of SEQ ID NO: 126 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 128;
- (gg) a light chain variable region having the polypeptide sequence of SEQ ID NO: 130 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 132:
- (hh) a light chain variable region having the polypeptide sequence of SEQ ID NO: 134 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 136;
- (ii) a light chain variable region having the polypeptide sequence of SEQ ID NO: 138 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 140:
- (jj) a light chain variable region having the polypeptide sequence of SEQ ID NO: 142 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 144:
- (kk) a light chain variable region having the polypeptide sequence of SEQ ID NO: 146 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 148:
- (II) a light chain variable region having the polypeptide sequence of SEQ ID NO: 150 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 152:

- (mm) a light chain variable region having the polypeptide sequence of SEQ ID NO: 154 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 156:
- (nn) a light chain variable region having the polypeptide sequence of SEQ ID NO: 158 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 160:
- (oo) a light chain variable region having the polypeptide sequence of SEQ ID NO: 162 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 164.
- (pp) a light chain variable region having the polypeptide sequence of SEQ ID NO: 166 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 168:
- (qq) a light chain variable region having the polypeptide sequence of SEQ ID NO: 170 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 172; or
- (rr) a light chain variable region having the polypeptide sequence of SEQ ID NO: 86 and a heavy chain variable region having the polypeptide sequence of SEQ ID NO: 136.
- 9. An isolated monoclonal antibody that binds to human tissue factor pathway inhibitor, wherein the antibody comprises a human heavy chain variable region comprising an amino acid sequence having at least 96% identity to an amino acid sequence selected from the group consisting of the amino acid sequences set forth in SEQ ID NO:4, SEQ ID NO:8, SEQ ID NO:12, SEQ ID NO:16, SEQ ID NO:20, SEQ ID NO:24, SEQ ID NO:28, SEQ ID NO:32, SEQ ID NO:36, SEQ ID NO:40, SEQ ID NO:44, SEQ ID NO:48, SEQ ID NO:52, SEQ ID NO:56, SEQ ID NO:60, SEQ ID NO:64, SEQ ID NO:68, SEQ ID NO:72, SEQ ID NO:76, SEQ ID NO:80, SEO ID NO:84, SEO ID NO:88, SEO ID NO:92, SEQ ID NO:96, SEQ ID NO:100, SEQ ID NO:104, SEQ ID NO:108, SEQ ID NO:112, SEQ ID NO:116, SEQ ID NO:120, SEQ ID NO:124, SEQ ID NO:128, SEQ ID NO:132, SEQ ID NO:136, SEQ ID NO:140, SEQ ID NO:144, SEQ ID NO:148, SEQ ID NO:152, SEQ ID NO:156, SEQ ID NO:160, SEQ ID NO:164, SEQ ID NO:168, and SEQ ID NO:172.
- 10. An isolated monoclonal antibody that binds to human tissue factor pathway inhibitor, wherein the antibody comprises a human light chain variable region comprising an amino acid sequence having at least 97% identity to an amino acid sequence selected from the group consisting of the amino acid sequences set forth in SEQ ID NO:2, SEQ ID NO:6, SEQ ID NO:10, SEQ ID NO:14, SEQ ID NO:18, SEQ ID NO:22, SEQ ID NO:26, SEQ ID NO:30, SEQ ID NO:34, SEQ ID NO:38, SEQ ID NO:42, SEQ ID NO:46, SEQ ID NO:50, SEQ ID NO:54, SEQ ID NO:58, SEQ ID NO:62, SEQ ID NO:66, SEQ ID NO:70, SEQ ID NO:74, SEQ ID NO:78, SEQ ID NO:82, SEQ ID NO:86, SEQ ID NO:90, SEQ ID NO:94, SEQ ID NO:98, SEQ ID NO:102, SEQ ID NO:106, SEQ ID NO:110, SEQ ID NO:114, SEQ ID NO:118, SEQ ID NO:122, SEQ ID NO:126, SEQ ID NO:130, SEQ ID NO:134, SEQ ID NO:138, SEQ ID NO:142, SEQ ID NO:146, SEQ ID NO:150, SEQ ID NO:154, SEQ ID NO:158, SEQ ID NO:162, SEQ ID NO:166, and SEQ ID NO:170.
- 11. The antibody of claim 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10, wherein the antibody is selected from the group consisting of

- an IgG1, an IgG2, an IgG3, an IgG4, an IgM, an IgA1, an IgA2, a secretory IgA, an IgD, and an IgE antibody.
- 12. The antibody of claim 1, 2, 3, 4, 5, 6, 7, 8, 9, 10 or 11, wherein blood clotting time in the presence of the antibody is shortened as measured by diluted prothrombin time.
- 13. The antibody of claim 1, 2, 3, 4, 5, 6, 7, 8, 9, 10 or 11 that is an antibody fragment or a single chain antibody.
- 14. A pharmaceutical composition comprising a therapeutically effective amount of the monoclonal antibody of claim 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12 or 13 and a pharmaceutically acceptable carrier.
- 15. A method for treating genetic and acquired deficiencies or defects in coagulation comprising administering a therapeutically effective amount of the pharmaceutical composition of claim 14 to a patient.
- **16**. The method of claim **15**, wherein the method treats hemophilia A or B.
- 17. A method for shortening bleeding time comprising administering a therapeutically effective amount of the pharmaceutical composition of claim 14 to a patient.
- 18. A method for treating genetic and acquired deficiencies in coagulation comprising administering (a) a first amount of a monoclonal antibody that binds to human tissue factor pathway inhibitor and (b) a second amount of factor VIII or factor IX, wherein said first and second amounts together are effective for treating said deficiencies or defects, and further wherein factor VII is not coadministered.
- 19. A pharmaceutical composition comprising a therapeutically effective amount of the combination of (a) a monoclonal antibody of claim 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12 or 13 and (b) factor VIII or factor IX; wherein the composition does not contain factor VII.
- 20. A method for treating genetic and acquired deficiencies or defects in coagulation comprising administering a therapeutically effective amount of the pharmaceutical composition of claim 19 to a patient in need thereof.
- 21. A method for shortening bleeding time comprising administering a therapeutically effective amount of the pharmaceutical composition of claim 19 to a patient.
- **22**. An isolated fully human monoclonal antibody to human tissue factor pathway inhibitor.
- 23. A pharmaceutical composition comprising a therapeutically effective amount of the monoclonal antibody of claim 22 and a pharmaceutically acceptable carrier.
- 24. A method for treating genetic and acquired deficiencies or defects in coagulation comprising administering a therapeutically effective amount of the pharmaceutical composition of claim 23 to a patient.
- **25**. The method of claim **24**, further comprising administering with factor VIII or factor IX.

- 26. An isolated nucleic acid molecule encoding an antibody that binds to human tissue factor pathway inhibitor, wherein the antibody comprises a CDR3 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 388-430.
- 27. An isolated nucleic acid molecule encoding an antibody that binds to human tissue factor pathway inhibitor, wherein the antibody comprises a CDR3 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 259-301.
- **28**. A method for producing a fully human monoclonal antibody that binds human tissue factor pathway inhibitor comprising:
  - (i) transfecting a nucleotide sequence encoding the fully human monoclonal antibody into a host cell, and
  - (ii) culturing the host cell so to express the monoclonal antibody.
- 29. The method of claim 28, wherein the monoclonal antibody comprises a CDR3 comprising an amino acid sequence selected from the group of sequences consisting of SEQ ID NOs: 388-430
- **30**. A method for producing a monoclonal antibody that binds human tissue factor pathway inhibitor comprising:
  - (i) transfecting a nucleotide sequence encoding the monoclonal antibody into a host cell, and
  - (ii) culturing the host cell so to express the monoclonal antibody,
  - wherein the monoclonal antibody comprises a CDR3 comprising an amino acid sequence selected from the group of sequences consisting of SEQ ID NOs: 259-301.
- 31. The method of claim 29, wherein the monoclonal antibody comprises a heavy chain CDR3 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 388-430 and a light chain CDR3 comprising an amino amino acid sequence selected from the group of sequences consisting of SEQ ID NOs: 259-301.
- **32.** An isolated fully human monoclonal antibody to Kunitz domain 2 of human tissue factor pathway inhibitor.
- **33**. A pharmaceutical composition comprising a therapeutically effective amount of the monoclonal antibody of claim **32** and a pharmaceutically acceptable carrier.
- **34**. A method for treating genetic and acquired deficiencies or defects in coagulation comprising administering a therapeutically effective amount of the pharmaceutical composition of claim **33** to a patient.
- **35**. The method of claim **34**, further comprising administering with factor VII, factor VIII or factor IX.

\* \* \* \* \*