

February 2007

# Supplementary Data for “Sequence and annotation of the 314-kb MT325 and the 321-kb FR483 viruses that infect *Chlorella Pbi*”: Appendix D: Gene Names N003L through N847R

Lisa A. Fitzgerald  
*University of Nebraska-Lincoln, fitz918@msn.com*

Michael V. Graves  
*University of Massachusetts-Lowell, Michael\_Graves@uml.edu*

Xiao Li  
*University of Massachusetts-Lowell*

Tamara Feldblyum  
*The Institute for Genomic Research, Rockville, MD*

James Hartigan  
*Agencourt Bioscience Corporation, Beverly, MA*

*See next page for additional authors*

Follow this and additional works at: <http://digitalcommons.unl.edu/virologypub>

 Part of the [Virology Commons](#)

---

Fitzgerald, Lisa A.; Graves, Michael V.; Li, Xiao; Feldblyum, Tamara; Hartigan, James; and Van Etten, James L., "Supplementary Data for “Sequence and annotation of the 314-kb MT325 and the 321-kb FR483 viruses that infect *Chlorella Pbi*”: Appendix D: Gene Names N003L through N847R” (2007). *Virology Papers*. 5.  
<http://digitalcommons.unl.edu/virologypub/5>

This Article is brought to you for free and open access by the Virology, Nebraska Center for at DigitalCommons@University of Nebraska - Lincoln. It has been accepted for inclusion in Virology Papers by an authorized administrator of DigitalCommons@University of Nebraska - Lincoln.

---

**Authors**

Lisa A. Fitzgerald, Michael V. Graves, Xiao Li, Tamara Feldblyum, James Hartigan, and James L. Van Etten

SUPPLEMENTARY DATA FOR

# Sequence and annotation of the 314-kb MT325 and the 321-kb FR483 viruses that infect *Chlorella Pbi*

Lisa A. Fitzgerald<sup>a</sup>, Michael V. Graves<sup>b</sup>, Xiao Li<sup>b</sup>, Tamara Feldblyum<sup>c</sup>, James Hartigan<sup>d</sup>, and James L. Van Etten<sup>e, f, \*</sup>

<sup>a</sup>Department of Chemistry, University of Nebraska–Lincoln, Lincoln, NE 68588-0304

<sup>b</sup>Department of Biological Sciences, University of Massachusetts–Lowell, Lowell, MA 01854

<sup>c</sup>The Institute for Genomic Research, 9712 Medical Center Drive, Rockville, MD 20850

<sup>d</sup>Agencourt Bioscience Corporation, 500 Cummings Center, Suite 2450, Beverly, MA 01915

<sup>e</sup>Department of Plant Pathology, University of Nebraska–Lincoln, Lincoln, NE 68583-0722

<sup>f</sup>Nebraska Center for Virology, University of Nebraska, Lincoln, NE 68588-0666

\*Corresponding author. Email: jvanetten@unlnotes.unl.edu

**Abstract:** Viruses MT325 and FR483, members of the family Phycodnaviridae, genus *Chlorovirus*, infect the fresh water, unicellular, eukaryotic, chlorella-like green alga, *Chlorella Pbi*. The 314,335-bp genome of MT325 and the 321,240-bp genome of FR483 are the first viruses that infect *Chlorella Pbi* to have their genomes sequenced and annotated. Furthermore, these genomes are the two smallest chlorella virus genomes sequenced to date, MT325 has 331 putative protein-encoding and 10 tRNA-encoding genes and FR483 has 335 putative protein-encoding and 9 tRNA-encoding genes. The protein-encoding genes are almost evenly distributed on both strands, and intergenic space is minimal. Approximately 40% of the viral gene products resemble entries in public databases, including some that are the first of their kind to be detected in a virus. For example, these unique gene products include an aquaglyceroporin in MT325, a potassium ion transporter protein and an alkyl sulfatase in FR483, and a dTDP–glucose pyrophosphorylase in both viruses. Comparison of MT325 and FR483 protein-encoding genes with the prototype chlorella virus PBCV-1 indicates that approximately 82% of the genes are present in all three viruses.

Supplementary data associated with this article is archived in this repository as 4 separate files: Appendices A–D. Each document, in spreadsheet format, shows Gene Name, Genome Position, A.A. length, Peptide Mw, pI, CDD Hit Number, COGs, COG Definition, Bit Score, E-value, % Identity, % Positive, Query from-to, Hit from-to, BLASTp Hit Number, Hit Accession, BLASTp Definition, Bit Score, E-value, % Identity, % Positive, Query from-to, and Hit from-to.

Appendix A: Gene Names m002R through m843L

Appendix B: Gene Names M001L through M807R

Appendix C: Gene Names n001L through n849R

Appendix D: Gene Names N003L through N847R

---

## Appendix D: Gene Names N003L through N847R

Gene Name	Genome Position	A.A. length	Peptide Mw	pI	CDD Hit Number	COGs	COG Definition	Bit Score	E-value	% Identity	% Positive	Query from-to	Hit from-to	BLASTp Hit Number	Hit Accession	BLASTp Definition	Bit Score	E-value	% Identity	% Positive	Query from-to	Hit from-to
N003L	2472-490	661	74,180	9.20	1	COG2015	COG2015, Alkyl sulfatase and related hydrolases [Secondary metabolites biosynthesis, transport, and catabolism]	700.55	0.00E+00	45%	64%	22-654	15-652	1	NP_896208	alkyl sulfatase family protein	752.67	0.00E+00	58%	73%	34-657	50-679
					2	pfam00753	Lactamase B, Metallo-beta-lactamase superfamily..	48.00	3.49E-06	22%	36%	129-232	1-102	2	AAZ27425	metallo-beta-lactamase family protein	741.50	0.00E+00	57%	74%	34-657	45-675
					3	COG1237	COG1237, Metal-dependent hydrolases of the beta-lactamase superfamily [General function prediction only]	40.68	6.24E-04	27%	42%	138-224	26-107	3	CAG19676	hypothetical beta-lactamase	731.48	0.00E+00	56%	73%	34-658	48-678
					4	COG4783	COG4783, Putative Zn-dependent protease, contains TPR repeats [General function prediction only]	38.01	3.33E-03	33%	54%	464-531	378-445	4	AAV98272	metallo-beta-lactamase family protein	721.47	0.00E+00	56%	73%	34-657	34-663
					5	COG0491	Glob, Zn-dependent hydrolases, including glyoxylases [General function prediction only]	35.99	1.66E-02	29%	44%	138-223	30-109	5	YP_593031	beta-lactamase-like	671.77	0.00E+00	52%	70%	34-658	38-667
					6	COG1234	ElAC, Metal-dependent hydrolases of the beta-lactamase superfamily III [General function prediction only]	35.80	1.93E-02	24%	48%	163-215	44-98	6	ZP_00753431	COG2015: Alkyl sulfatase and related hydrolases	620.93	4.82E-176	48%	67%	30-658	26-660
					7	cd00189	TPR, Tetratricopeptide repeat domain; typically contains 34 amino acids [WLP]-X(2)-[LIM]-[GAS]-X(2)-[YLF]-X(8)-[ASE]-X(3)-[FPL]-X(2)-[ASL]-X(4)-[PKE] is the consensus sequence; found in a variety of organisms including bacteria, cyanobacteria, yeast, fungi, plants, and humans in various subcellular locations; involved in a variety of functions including protein-protein interactions, but common features in the interaction partners have not been defined; involved in chaperone, cell-cycle, transcription, and protein transport complexes; the number of TPR motifs varies among proteins (1,3-11,13,15,16,19); 5-6 tandem repeats generate a right-handed helical structure with an amphipathic channel that is thought to accommodate an alpha-helix of a target protein; it has been proposed that TPR proteins preferably interact with WD-40 repeat proteins, but in many instances several TPR-proteins seem to aggregate to multi-protein complexes; examples of TPR-proteins include, Cdc16, Cdc-23n and Cdc-27n components of the ribosomal/APC superfamily	33.90	7.00E-02	34%	50%	464-508	4-48	7	ZP_00750973	COG2015: Alkyl sulfatase and related hydrolases	620.54	6.29E-176	48%	67%	30-658	26-660
					8	COG2333	ComEC, Predicted hydrolase (metallo-beta-lactamase superfamily) [General function prediction only]	32.23	2.16E-01	25%	43%	137-224	57-138	8	ZP_00747297	COG2015: Alkyl sulfatase and related hydrolases	619.77	1.07E-175	48%	67%	31-658	27-660
					9	COG2813	RsmC, 16S RNA G1207 methylase RsmC [Translation, ribosomal structure and biogenesis]	31.39	3.98E-01	23%	45%	508-600	126-212	9	AAF93948	conserved hypothetical protein	619.00	1.83E-175	48%	67%	31-658	27-660
					10	COG0595	COG0595, Predicted hydrolase of the metallo-beta-lactamase superfamily [General function prediction only]	30.60	5.81E-01	23%	42%	135-189	23-83	10	ZP_00913757	Twin-arginine translocation pathway signal	605.52	2.09E-171	49%	66%	34-657	49-678
N007L	3122-2577	182	20,422	10.22		No Hit Found								1	AAC96449	A81L	108.61	9.58E-23	41%	62%	45-158	37-163
N010L	3666-3151	172	19,292	6.65		No Hit Found								1	AAC96452	A84L	71.63	1.10E-11	32%	55%	14-135	13-149
N012L	4563-3751	271	31,655	6.29	1	pfam03016	Exostosin, Exostosin family. The EXT family is a family of tumour suppressor genes. Mutations of EXT1 on 8q24, EXT2 on 11p11-13, and EXT3 on 19p have been associated with the autosomal dominant disorder known as hereditary multiple exostoses (HME). This is the most common known skeletal dysplasia. The chromosomal locations of other EXT genes suggest association with other forms of neoplasia. EXT1 and EXT2 have both been shown to encode a heparan sulphate polymerase with both D-glucuronyl (GlcA) and N-acetyl-D-glucosaminoyl (GlcNAc) transferase activities. The nature of the defect in heparan sulphate biosynthesis in HME is unclear.	45.82	6.04E-06	25%	51%	159-235	221-292	1	AAC96443	A75L	234.57	2.74E-60	41%	61%	8-268	7-276
					2	pfam01323	DSBA, DSBA-like thioesteron domain. This family contains a diverse set of proteins with a thioesteron-like structure pfam0095. This family also includes 2-hydroxychromene-2-carboxylate (HCCA) isomerase enzymes catalyse one step in prokaryotic polyaromatic hydrocarbon (PAH) catabolic pathways. This family also contains members with functions other than HCCA isomerisation, such as Kappa family GSTs, whose similarity to HCCA isomerases was not previously recognised. Some members of this family may have been mis-annotated in protein sequence databases.	29.98	3.06E-01	17%	34%	43-161	8-138	2	NP_195517	catalytic	45.82	1.80E-03	27%	48%	162-268	284-384
					3	NP_195005	catalytic	44.67	4.02E-03	26%	49%	174-266	471-571									
					4	BAC42936	unknown protein	44.67	4.02E-03	26%	49%	174-266	148-248									
					5	ABA18110	exostosin family protein	43.51	8.95E-03	30%	44%	169-257	206-308									
N014L	5025-4663	121	13,198	10.46	1	cd01285	nucleoside_deaminase, Nucleoside deaminases include adenosine, guanine and cytosine deaminases. These enzymes are Zn dependent and catalyze the deamination of nucleosides. The zinc ion in the active site plays a central role in the proposed catalytic mechanism, activating a water molecule to form a hydroxide ion that performs a nucleophilic attack on the substrate. The functional enzyme is a homodimer. Cytosine deaminase catalyzes the deamination of cytosine to uracil and ammonia and is a member of the pyrimidine salvage pathway. Cytosine deaminase is found in bacteria and fungi but is not present in mammals; for this reason, the enzyme is currently of interest for antimicrobial drug design and gene therapy applications against tumors. Some members of this family are tRNA-specific adenosine deaminases that generate inosine at the first position of their anticodon (position 34) of specific tRNAs; this modification is thought to enlarge the codon recognition capacity during protein synthesis. Other members of the family are guanine deaminases which deaminate guanine to xanthine as part of the utilization of guanine.	58.35	2.32E-10	37%	51%	4-106	1-94	1	AAC96568	contains cytidine and deoxycytidine deaminase Zn-binding region signature	159.84	2.40E-38	61%	83%	1-118	1-118
					2	COG0590	CumB, Cytosine/adenosine deaminases [Nucleotide transport and metabolism / Translation, ribosomal structure and biogenesis]	53.42	6.92E-09	33%	51%	2-106	10-105	2	AAR26853	FirV-1-A29	50.83	1.57E-05	27%	50%	22-111	24-105
					3	pfam00383	dCMP_cyt_deam, Cytidine and deoxycytidylate deaminase zinc-binding region.	44.20	4.27E-06	31%	47%	4-106	7-101	3	AAV51127	tRNA-specific adenosine deaminase	50.06	2.67E-05	32%	49%	4-108	10-104
					4	cd01284	Riboflavin_deaminase-reductase, Riboflavin-specific deaminase. Riboflavin biosynthesis protein RbD (Diaminohydroxyphosphoribosylamine deaminase) catalyzes the deamination of 2,5-diamino-6-ribosylamino-(4H)-pyrimidinone 5&apos;phosphate, which is an intermediate step in the biosynthesis of riboflavin. The rbg gene of Bacillus subtilis and the rbd gene of E. coli are bifunctional and contain this deaminase domain and a reductase domain which catalyzes the subsequent reduction of the ribosyl side chain.	37.92	3.09E-04	27%	45%	4-107	1-95	4	AAC68441	cytosine deaminase	49.68	3.49E-05	32%	49%	4-108	10-104
					5	cd01286	deoxycytidylate_deaminase, Deoxycytidylate deaminase domain. Deoxycytidylate deaminase catalyzes the deamination of dCMP to dUMP, providing the nucleotide substrate for thymidylate synthase. The enzyme binds Zn++, which is required for catalytic activity. The activity of the enzyme is allosterically regulated by the ratio of dCTP to dTTP not only in eukaryotic cells but also in T-even phage-infected Escherichia coli, with dCTP acting as an activator and dTTP as an inhibitor.	31.78	2.25E-02	37%	51%	50-108	68-117	5	ZP_00679087	Cytidine/deoxycytidylate deaminase, zinc-binding region	45.44	6.58E-04	32%	44%	4-106	19-111
					6	COG0117	RbD, Pyrimidine deaminase [Coenzyme metabolism].	31.76	2.38E-02	24%	43%	7-109	13-104	6	GAA00608	unnamed protein product	44.67	1.12E-03	30%	46%	4-108	36-130
					7	cd00786	cytidine_deaminase-like, Cytidine and deoxycytidylate deaminase zinc-binding region. The family contains cytidine deaminases, nucleoside deaminases, deoxycytidylate deaminases and riboflavin deaminases. Also included are the specific family of tRNA editing enzymes. All members are Zn dependent. The zinc ion in the active site plays a central role in the proposed catalytic mechanism, activating a water molecule to form a hydroxide ion that performs a nucleophilic attack on the substrate.	30.37	5.89E-02	20%	36%	6-108	3-95	7	NP_502546	JC8.4	44.28	1.47E-03	32%	48%	2-108	11-111
					8	pfam02041	Auxin BP, Auxin binding protein..	28.08	3.02E-01	21%	50%	23-79	58-126	8	AAF73539	cytidine/deoxycytidylate deaminase family protein	44.28	1.47E-03	29%	46%	4-108	10-104
					9	COG2131	ComEB, Deoxycytidylate deaminase [Nucleotide transport and metabolism].	26.85	6.80E-01	45%	60%	88-108	106-126	9	ABB14795	cytidine/deoxycytidylate deaminase family protein	43.90	1.92E-03	29%	46%	6-108	9-101
10	YP_428908	CMP/dCMP deaminase, zinc-binding	43.51	2.50E-03	31%	43%	6-108	9-101														
N015L	5407-5111	99	10,616	10.54		No Hit Found								1	AAC96567	A199R	70.86	1.48E-11	47%	66%	1-71	1-72
N016R	5419-5907	163	18,543	8.04		No Hit Found								1	AAC96564	A196L	184.88	7.72E-46	61%	79%	23-156	18-151

Gene Name	Genome Position	A.A. Length	Peptide Mw	pI	CDD Hit Number	COGs	COG Definition	Bit Score	E-value	% Identity	% Positive	Query from-to	Hit from-to	BLASTp Hit Number	Hit Accession	BLASTp Definition	Bit Score	E-value	% Identity	% Positive	Query from-to	Hit from-to
N019R	5958-6755	266	29,728	5.00	1	cd00577	PCNA, Proliferating Cell Nuclear Antigen (PCNA) domain found in eukaryotes and archaea. These polymerase processivity factors play a role in DNA replication and repair. PCNA encodes duplex DNA in its central cavity, providing a DNA-bound platform for the attachment of the polymerase. The trimeric PCNA ring is structurally similar to the dimeric ring formed by the DNA polymerase processivity factors in bacteria (beta subunit DNA polymerase III holoenzyme) and in bacteriophages (galactylic subunits in T4 and RB69). This structural correspondence further substantiates the mechanistic connection between eukaryotic and prokaryotic DNA replication that has been suggested on biochemical grounds. PCNA is also involved with proteins involved in cell cycle processes such as DNA repair and apoptosis. Many of these proteins contain a highly conserved motif known as the PIP-box (PCNA interacting protein box) which contains the sequence Qxx[LIM]xx[FY].	146.96	2.13E-36	30%	55%	12-261	1-245	1	AAC96561	similar to human PCNA, corresponds to Swis-Prot Accession Number P12004	368.24	1.53E-100	68%	85%	9-261	7-259
					2	pfam00705	PCNA_N, Proliferating cell nuclear antigen, N-terminal domain. N-terminal and C-terminal domains of PCNA are topologically identical. Three PCNA molecules are tightly associated to form a closed ring encircling duplex DNA.	92.25	5.33E-20	28%	57%	9-132	1-125	2	XP_534355	PREDICTED: similar to proliferating cell nuclear antigen	158.69	1.84E-37	32%	55%	5-264	202-461
					3	pfam02747	PCNA_C, Proliferating cell nuclear antigen, C-terminal domain. N-terminal and C-terminal domains of PCNA are topologically identical. Three PCNA molecules are tightly associated to form a closed ring encircling duplex DNA.	79.98	2.97E-16	34%	55%	137-261	2-127	3	CAA55669	proliferative cell nuclear antigen	157.53	4.10E-37	34%	56%	9-264	1-256
					4	COG0592	DnaN, DNA polymerase sliding clamp subunit (PCNA homolog) [DNA reinitiation, recombination, and repair]	56.08	4.36E-09	19%	37%	21-261	72-321	4	AAB27811	PCNA	156.76	6.99E-37	32%	57%	9-264	1-256
					5	COG1092	COG1092, Predicted SAM-dependent methyltransferases [DNA function prediction onlv].	31.44	1.30E-01	20%	31%	106-202	251-358	5	NP_172217	PCNA1 (PROLIFERATING CELLULAR NUCLEAR ANTIGEN); DNA binding / DNA polymerase processivity factor	156.76	6.99E-37	32%	57%	9-264	1-256
					6								6	AAG10077	proliferating cell nuclear antigen	155.99	1.10E-36	32%	55%	9-264	1-256	
					7								7	CAA37243	unnamed protein product	155.61	1.56E-36	32%	55%	9-264	1-256	
					8								8	BAB28355	unnamed protein product	155.61	1.56E-36	32%	55%	9-264	1-256	
					9								9	NP_035175	proliferating cell nuclear antigen	155.61	1.56E-36	32%	55%	9-264	1-256	
					10								10	NP_001029666	hypothetical protein LOC515499	155.22	2.04E-36	32%	55%	9-264	1-256	
N020L	10628-6753	1292	141,764	11.55	1	cd03274	ABC_SMC4_euk, Eukaryotic SMC4 proteins; SMC proteins are large (approximately 110 to 170 kDa), and each is arranged into five recognizable domains. Amino-acid sequence homology of SMC proteins between species is largely confined to the amino- and carboxy-terminal globular domains. The amino-terminal domain contains a &Apos;Walker A&apos; nucleotide-binding domain (GxxGxGKS/T, in the single-letter amino-acid code), which by mutational studies has been shown to be essential in several proteins. The carboxy-terminal domain contains a sequence (the DA-box) that resembles a &Apos;Walker B&apos; motif, and a motif with homology to the signature sequence of the ATP-binding cassette (ABC) family of ATPases. The sequence homology within the carboxy-terminal domain is relatively high within the SMC1-SMC4 group, whereas SMC5 and SMC6 show some divergence in both of these sequences. In eukaryotic cells, the proteins are found as heterodimers of SMC1 paired with SMC3, SMC2 with SMC4, and SMC5 with SMC6 (formerly known as Rad18).	33.04	2.36E-01	17%	51%	1073-1156	287-376	1	AAC96557	similar to SWI/SNF chromatin remodeling complex subunit OSA2	718.77	0.00E+00	39%	51%	2-1175	10-1139
					2	pfam01576	Myosin_tail_1, Myosin tail. The myosin molecule is a multi-subunit complex made up of two heavy chains and four light chains. It is a fundamental contractile protein found in all eukaryotic cell types. This family consists of the coiled-coil myosin heavy chain tail region. The coiled-coil is composed of the tail from two molecules of myosin. These can then assemble into the macromolecular thick filament. The coiled-coil region provides the structural backbone of the thick filament.	31.82	5.49E-01	34%	55%	1073-1155	772-846	2	T17682	hypothetical protein A192R - Chlorella virus PBCV-1	81.65	2.26E-13	43%	68%	1075-1175	103-200
					3	cd03275	ABC_SMC1_euk, Eukaryotic SMC1 proteins; SMC proteins are large (approximately 110 to 170 kDa), and each is arranged into five recognizable domains. Amino-acid sequence homology of SMC proteins between species is largely confined to the amino- and carboxy-terminal globular domains. The amino-terminal domain contains a &Apos;Walker A&apos; nucleotide-binding domain (GxxGxGKS/T, in the single-letter amino-acid code), which by mutational studies has been shown to be essential in several proteins. The carboxy-terminal domain contains a sequence (the DA-box) that resembles a &Apos;Walker B&apos; motif, and a motif with homology to the signature sequence of the ATP-binding cassette (ABC) family of ATPases. The sequence homology within the carboxy-terminal domain is relatively high within the SMC1-SMC4 group, whereas SMC5 and SMC6 show some divergence in both of these sequences. In eukaryotic cells, the proteins are found as heterodimers of SMC1 paired with SMC3, SMC2 with SMC4, and SMC5 with SMC6 (formerly known as Rad18).	31.95	5.73E-01	19%	44%	1063-1170	245-353	3	AAK77699	ORF30, putative collagen	50.83	4.27E-04	23%	28%	98-482	666-1074
					4	pfam06519	ToIA, ToIA protein. This family consists of several bacterial ToIA proteins as well as two eukaryotic proteins of unknown function. ToI proteins are involved in the translocation of group A colicins. Colicins are bacterial protein toxins, which are active against Escherichia coli and other related species (See pfam01024). ToIA is anchored to the cytoplasmic membrane by a single membrane spanning segment near the N-terminus, leaving most of the protein exposed to the periplasm.	31.64	6.12E-01	29%	45%	1070-1168	67-171	4	NP_477523	wsv001	50.83	4.27E-04	23%	28%	98-482	666-1074
					5	cd03276	ABC_SMC6_euk, Eukaryotic SMC6 proteins; SMC proteins are large (approximately 110 to 170 kDa), and each is arranged into five recognizable domains. Amino-acid sequence homology of SMC proteins between species is largely confined to the amino- and carboxy-terminal globular domains. The amino-terminal domain contains a &Apos;Walker A&apos; nucleotide-binding domain (GxxGxGKS/T, in the single-letter amino-acid code), which by mutational studies has been shown to be essential in several proteins. The carboxy-terminal domain contains a sequence (the DA-box) that resembles a &Apos;Walker B&apos; motif, and a motif with homology to the signature sequence of the ATP-binding cassette (ABC) family of ATPases. The sequence homology within the carboxy-terminal domain is relatively high within the SMC1-SMC4 group, whereas SMC5 and SMC6 show some divergence in both of these sequences. In eukaryotic cells, the proteins are found as heterodimers of SMC1 paired with SMC3, SMC2 with SMC4, and SMC5 with SMC6 (formerly known as Rad18).	31.16	8.12E-01	19%	43%	1073-1169	262-364	5	T17681	hypothetical protein a191R - Chlorella virus PBCV-1	59.31	1.20E-06	46%	65%	843-911	3-63
													6	NP_001015110	CG40323-PB.3	48.91	1.62E-03	23%	31%	96-472	3426-3778	
													7	XP_783728	PREDICTED: similar to Protein transport protein Sec24C (SEC24-related protein C)	55.07	2.26E-05	25%	33%	190-498	6-311	
													8	NP_571069	calymin	52.37	1.47E-04	23%	32%	140-483	230-604	
													9	AAH90694	Crm protein	52.37	1.47E-04	23%	32%	140-483	222-596	
													10	XP_394285	PREDICTED: similar to GA11046-PA	53.14	8.60E-05	27%	35%	192-481	124-394	
N023L	13471-10658	938	106,276	7.86	1	COG0417	PolB, DNA polymerase elongation subunit (family B) [DNA replication, recombination, and repair].	346.65	6.28E-96	30%	48%	51-879	11-772	1	AAC96553	PBVC-1 DNA polymerase	1352.04	0.00E+00	72%	84%	30-932	1-913
					2	pfam00136	DNA_pol_B, DNA polymerase family B. This region of DNA polymerase B appears to consist of more than one structural domain, possibly including elongation, DNA-binding and dNTP-binding activities.	337.35	4.53E-93	38%	55%	456-879	1-439	2	BAA35142	DNA polymerase	1347.03	0.00E+00	72%	83%	30-932	1-913
					3	smart00486	FOLBc, DNA polymerase type-B family. DNA polymerase alpha, delta, epsilon and zeta chain (eukaryota), DNA polymerases in archaea, DNA polymerase II in e. coli, mitochondrial DNA polymerases and and virus DNA polymerases.	292.12	1.89E-79	33%	51%	205-667	1-475	3	P30320	DNA polymerase	1341.25	0.00E+00	71%	83%	30-932	1-913
					4	cd00145	FOLBc, DNA polymerase type-B family. DNA directed DNA polymerase. Possesses DNA binding, polymerase and 3&apos;5&apos;exonuclease activity.	270.64	5.27E-73	33%	49%	205-704	1-511	4	AAB49748	DNA polymerase	443.74	1.58E-122	96%	97%	489-718	1-230
					5	pfam03104	DNA_pol_B_exo, DNA polymerase family B, exonuclease domain. This domain has 3&apos;5&apos;exonuclease activity and adopts a ribonuclease H type fold.	194.94	3.39E-50	26%	41%	56-383	1-334	5	AAK28935	DNA polymerase	431.80	6.23E-119	100%	100%	497-711	1-215
													6	AAK28933	DNA polymerase	429.10	4.04E-118	99%	100%	497-711	1-215	
													7	AAK86472	DNA polymerase	429.10	4.04E-118	99%	100%	498-711	1-214	











Gene Name	Genome Position	A.A. length	Peptide Mw	pI	CDD Hit Number	COGs	COG Definition	Bit Score	E-value	% Identity	% Positive	Query from-to	Hit from-to	BLASTp Hit Number	Hit Accession	BLASTp Definition	Bit Score	E-value	% Identity	% Positive	Query from-to	Hit from-to
					7	cd02183	GH16_GPI glucosyltransferase, GPI (glycosylphosphatidylinositol)-glucosyltransferase is a GPI-anchored membrane protein present in the fungal cell wall that is thought to play an important role in cell wall biosynthesis. GPI-glycosyltransferase belongs to a family of glycosyl hydrolases that includes lichenase, xyloglucan endotransglycosylase (XET), beta-agarase, kappa-carrageenase, endo-beta-1,3-glucanase, endo-beta-1,3-1,4-glucanase, and endo-beta-galactosidase, all of which have a conserved jelly roll fold with a deep active site channel harboring the catalytic residues. GH16_lichenase, Lichenase, also known as 1,3-1,4-beta-glucanase, is a member of glycosyl hydrolase family 16 that specifically cleaves 1,4-beta-D-glycosidic bonds in mixed-linked beta-glucans that also contain 1,3-beta-D-glycosidic linkages. Natural substrates of beta-glucanase are beta-glucans from grain endosperm cell walls or lichenan from the lichenist moss <i>Cetraria islandica</i> . This protein is found not only in bacteria but also in anaerobic fungi. This domain includes two seven-stranded antiparallel beta-sheets that are adjacent to one another forming a rimmed-invariant beta-sandwich structure. GH16_kappa-carrageenase, Kappa-carrageenase degrades kappa-carrageenans which are the gel-forming, sulfated 1,3-alpha-1,4-beta-galactans that make up the cell walls of marine red algae such as Rhodosphecia. Kappa-carrageenases exist in bacteria belonging to at least three phylogenetically distant branches, including pseudomonads, planctomycetes, and bacteroidetes. This domain adopts a curved beta-sandwich conformation, with a tunnel-shaped active site cavity referred to as a lichenin fold.	38.68	1.04E-03	28%	49%	190-283	58-144	7	YP_563606	glycoside hydrolase, family 16	111.69	4.06E-23	31%	44%	76-343	38-325
					8	cd02175	GH16_lichenase, Lichenase, also known as 1,3-1,4-beta-glucanase, is a member of glycosyl hydrolase family 16 that specifically cleaves 1,4-beta-D-glycosidic bonds in mixed-linked beta-glucans that also contain 1,3-beta-D-glycosidic linkages. Natural substrates of beta-glucanase are beta-glucans from grain endosperm cell walls or lichenan from the lichenist moss <i>Cetraria islandica</i> . This protein is found not only in bacteria but also in anaerobic fungi. This domain includes two seven-stranded antiparallel beta-sheets that are adjacent to one another forming a rimmed-invariant beta-sandwich structure. GH16_kappa-carrageenase, Kappa-carrageenase degrades kappa-carrageenans which are the gel-forming, sulfated 1,3-alpha-1,4-beta-galactans that make up the cell walls of marine red algae such as Rhodosphecia. Kappa-carrageenases exist in bacteria belonging to at least three phylogenetically distant branches, including pseudomonads, planctomycetes, and bacteroidetes. This domain adopts a curved beta-sandwich conformation, with a tunnel-shaped active site cavity referred to as a lichenin fold.	37.58	2.12E-03	29%	46%	126-281	30-157	8	ZP_00637494	Glucan endo-1,3-beta-D-glucosidase	109.00	2.63E-22	31%	43%	85-343	48-325
					9	cd02177	GH16_kappa-carrageenase, Kappa-carrageenase degrades kappa-carrageenans which are the gel-forming, sulfated 1,3-alpha-1,4-beta-galactans that make up the cell walls of marine red algae such as Rhodosphecia. Kappa-carrageenases exist in bacteria belonging to at least three phylogenetically distant branches, including pseudomonads, planctomycetes, and bacteroidetes. This domain adopts a curved beta-sandwich conformation, with a tunnel-shaped active site cavity referred to as a lichenin fold.	35.47	1.09E-02	30%	46%	81-212	3-117	9	NP_763201	Beta-glucanase/Beta-glucan synthetase	107.84	5.86E-22	28%	43%	49-343	12-333
					10	cd02176	GH16_XET, Xyloglucan endotransglycosylase (XETs) cleave and religate xyloglucan polymers in plant cell walls via a transglycosylation mechanism. Thus, XET is a key enzyme in all plant processes that require cell wall remodeling. Even though the overall structure of XET is a curved beta-sandwich similar to other enzymes in the glycosyl hydrolase family 16, parts of its substrate binding cleft are more reminiscent of the distantly related xylorin hwltranase family 7.	33.64	3.40E-02	34%	46%	190-283	50-142	10	BAD63242	endo-beta-1,3-glucanase	107.46	7.66E-22	31%	44%	83-343	31-277
N124R	60078-61757	560	63,740	8.49	1	COG1215	COG1215, Glycosyltransferases, probably involved in cell wall biosynthesis (Cell envelope biosynthesis, outer membrane). Chitin synthase 2, Chitin synthase. Members of this family are fungal chitin synthase EC2.4.1.16 enzymes. They catalyze chitin synthesis as follows: UDP-N-acetyl-D-glucosamine + ((1,4)-(N-acetyl-beta-D-glucosaminyl))N) &lt;-&gt; UDP + ((1,4)-(N-acetyl-beta-D-glucosaminyl))N+1).	90.76	4.57E-19	25%	44%	39-390	10-328	1	AAD26641	hyaluronan synthase	832.40	0.00E+00	69%	82%	1-550	8-557
					2	pfam03919	Glycosyl transferase. Diverse family, transferring sugar from UDP-glucose, UDP-N-acetyl galactosamine, GDP-mannose or GDP-abequose, to a range of substrates including cellulose, dolichol phosphate and teichoic acids.	54.92	2.35E-08	23%	44%	186-345	193-373	2	AAC84666	PBCV-1 hyaluronic acid synthetase	828.55	0.00E+00	69%	82%	1-550	8-557
					3	pfam00535	Glycosyl transferase. Diverse family, transferring sugar from UDP-glucose, UDP-N-acetyl galactosamine, GDP-mannose or GDP-abequose, to a range of substrates including cellulose, dolichol phosphate and teichoic acids.	47.14	5.08E-06	24%	43%	86-283	2-167	3	AAD26643	hyaluronan synthase	826.62	0.00E+00	66%	82%	1-550	8-557
					4								4	NP_037285	hyaluronan synthase 2	218.78	4.51E-55	29%	48%	7-530	12-532	
					5								5	AAI09072	hyaluronan synthase 2	218.39	5.89E-55	29%	48%	7-530	12-532	
					6								6	AAC53309	hyaluronan synthase 2	218.39	5.89E-55	29%	48%	7-530	12-532	
					7								7	BAC37753	unnamed protein product	218.39	5.89E-55	29%	48%	7-530	12-532	
					8								8	XP_528222	PREDICTED, similar to hyaluronan synthase 2	218.39	5.89E-55	29%	48%	7-530	24-544	
					9								9	BAB63264	hyaluronic acid synthase 2	216.85	1.71E-54	29%	48%	7-530	12-532	
					10								10	NP_999218	hyaluronan synthase 2	216.85	1.71E-54	29%	48%	7-530	12-532	
N128R	61944-62906	321	36,035	8.75	1	pfam01331	mRNA_cap_enzyme, mRNA capping enzyme, catalytic domain. This family represents the ATP binding catalytic domain of the mRNA capping enzyme.	97.66	1.64E-21	25%	43%	50-223	1-192	1	AAC96471	PBCV-1 mRNA guanylyltransferase	326.25	9.18E-88	52%	68%	2-314	12-324
					2	COG5226	CEC1, mRNA capping enzyme, guanylyltransferase (alpha) subunit (RNA processing and modification).	75.46	8.40E-15	21%	37%	20-317	17-362	2	1CKN_B	Chain B, Structure Of Guanylylated Mmna Capping Enzyme Complexed With Gtp	323.94	4.55E-87	52%	68%	2-314	12-324
					3	pfam03919	mRNA cap C, mRNA capping enzyme, C-terminal domain.	50.74	2.48E-07	26%	44%	228-315	3-110	3	AAL98788	putative RNA guanylyltransferase	71.25	5.32E-11	26%	42%	44-311	224-479
					4	COG1947	ispC, 4-diphosphoryl-D-2-methyl-D-erythritol 2-phosphate synthase (LidD metabolite).	29.47	5.92E-01	27%	38%	135-183	50-98	4	CAG09212	unnamed protein product	69.71	1.55E-10	24%	43%	9-317	234-560
					5								5	AAT68133	mRNA capping enzyme	68.55	3.45E-10	23%	44%	8-317	234-560	
					6								6	AA82373	mRNA capping enzyme	68.17	4.50E-10	24%	39%	44-311	220-478	
					7								7	NP_974263	mRNA guanylyltransferase/ phosphoprotein phosphatase/ protein tyrosine/serine/threonine phosphatase	67.78	5.88E-10	22%	41%	33-318	337-650	
					8								8	P78587	mRNA capping enzyme alpha subunit (mRNA guanylyltransferase) (GTP-RNA guanylyltransferase) (GTase)	65.47	2.92E-09	25%	39%	47-316	41-373	
					9								9	1P16_B	Chain B, Structure Of An Mmna Capping Enzyme Bound To The Phosphorylated Carboxyl-Terminal Domain Of Rna Polymerase Ii	63.93	8.50E-09	25%	38%	47-316	41-373	
					10								10	AAH67387	RNA guanylyltransferase and 5&apos;-phosphatase	62.39	2.47E-08	23%	42%	8-313	231-563	
N131L	63284-62922	121	13,791	7.42	1	COG5170	CDCS5, Serine/threonine protein phosphatase 2A, regulatory subunit (Signal transduction mechanisms).	26.99	6.38E-01	35%	54%	58-84	59-85	1	AAC96969	A645R	80.11	2.41E-14	34%	57%	1-117	1-121
N132L	64179-63336	281	31,701	5.23	1	pfam00443	UCH, Ubiquitin carboxyl-terminal hydrolase..	109.71	3.76E-25	20%	40%	4-275	6-312	1	AAC96473	contains ubiquitin carboxy-terminal hydrolase active sites; similar to human ubiquitin carboxyl-terminal hydrolase, corresponds to Swiss-Prot Accession Number Q09879	261.92	1.71E-68	45%	67%	1-277	1-280
					2	cd02257	Peptidase_C19, Peptidase C19 contains ubiquitinyl hydrolases. They are intracellular peptidases that remove ubiquitin molecules from polyubiquitinated peptides by cleavage of isopeptide bonds. They hydrolyze bonds involving the carboxyl group of the C-terminal Gly residue of ubiquitin. The purpose of the de-ubiquitination is thought to be editing of the ubiquitin conjugates, which could rescue them from degradation, as well as recycling of the ubiquitin. The ubiquitin-proteasome system is responsible for most protein turnover in the mammalian cell, and with over 50 members, family C19 is one of the largest families of peptidases in the human genome.	81.90	8.61E-17	17%	33%	4-276	2-320	2	XP_654739	ubiquitin carboxyl-terminal hydrolase	60.46	7.57E-08	23%	41%	4-278	68-347
					3	cd02674	Peptidase_C19R, A subfamily of peptidase C19, Peptidase C19 contains ubiquitinyl hydrolases. They are intracellular peptidases that remove ubiquitin molecules from polyubiquitinated peptides by cleavage of isopeptide bonds. They hydrolyze bonds involving the carboxyl group of the C-terminal Gly residue of ubiquitin. The purpose of the de-ubiquitination is thought to be editing of the ubiquitin conjugates, which could rescue them from degradation, as well as recycling of the ubiquitin. The ubiquitin-proteasome system is responsible for most protein turnover in the mammalian cell, and with over 50 members, family C19 is one of the largest families of peptidases in the human genome.	74.61	1.15E-14	21%	35%	6-276	4-335	3	EA096319	Ubiquitin carboxyl-terminal hydrolase family protein	55.07	3.18E-06	20%	42%	6-277	425-736
					4	cd02661	Peptidase_C19E, A subfamily of Peptidase C19, Peptidase C19 contains ubiquitinyl hydrolases. They are intracellular peptidases that remove ubiquitin molecules from polyubiquitinated peptides by cleavage of isopeptide bonds. They hydrolyze bonds involving the carboxyl group of the C-terminal Gly residue of ubiquitin. The purpose of the de-ubiquitination is thought to be editing of the ubiquitin conjugates, which could rescue them from degradation, as well as recycling of the ubiquitin. The ubiquitin-proteasome system is responsible for most protein turnover in the mammalian cell, and with over 50 members, family C19 is one of the largest families of peptidases in the human genome.	63.74	2.55E-11	19%	37%	5-275	5-303	4	NP_990257	ubiquitin specific protease 2	52.76	1.58E-05	23%	39%	6-275	23-348
					5	cd02663	Peptidase_C19G, A subfamily of Peptidase C19, Peptidase C19 contains ubiquitinyl hydrolases. They are intracellular peptidases that remove ubiquitin molecules from polyubiquitinated peptides by cleavage of isopeptide bonds. They hydrolyze bonds involving the carboxyl group of the C-terminal Gly residue of ubiquitin. The purpose of the de-ubiquitination is thought to be editing of the ubiquitin conjugates, which could rescue them from degradation, as well as recycling of the ubiquitin. The ubiquitin-proteasome system is responsible for most protein turnover in the mammalian cell, and with over 50 members, family C19 is one of the largest families of peptidases in the human genome.	44.15	2.15E-05	37%	63%	213-261	278-332	5	EAS01964	Ubiquitin carboxyl-terminal hydrolase family protein	51.60	3.52E-05	21%	38%	6-259	186-497







Gene Name	Genome Position	A.A. length	Peptide Mw	pI	CDD Hit Number	COGs	COG Definition	Bit Score	E-value	% Identity	% Positive	Query from-to	Hit from-to	BLASTp Hit Number	Hit Accession	BLASTp Definition	Bit Score	E-value	% Identity	% Positive	Query from-to	Hit from-to
					5	pfam00270	DEAD, DEAD/DEAH box helicase. Members of this family include the DEAD and DEAH box helicases. Helicases are involved in unwinding nucleic acids. The DEAD box helicases are involved in various aspects of RNA metabolism, including nuclear transcription, pre mRNA splicing, ribosome biogenesis, nucleocytoplasmic transport, translation, RNA decay and noncellular gene expression	44.64	2.48E-05	18%	32%	102-233	19-175	5	YP_437404	DNA or RNA helicase of superfamily II	140.58	1.20E-31	29%	46%	87-430	421-776
					6	COG4096	HsdR, Type I site-specific restriction-modification system, R (restriction) subunit and related helicases (Defense mechanisms)	43.81	4.13E-05	26%	40%	113-231	190-321	6	YP_01111109	DNA or RNA helicase of superfamily II	139.81	2.04E-31	29%	46%	85-443	419-793
					7	pfam00176	SNF2_N, SNF2 family N-terminal domain. This domain is found in proteins involved in a variety of processes including transcription regulation (e.g., SNF2, STH1, brahma, MOT1), DNA repair (e.g., ERCC2, RAD18, RAD51), DNA recombination (e.g., RAD54) and chromatin unwinding (e.g., ISWI) as well as a variety of other proteins with little functional information (e.g. Invictar FTI 1)	43.73	5.08E-05	22%	38%	110-230	18-156	7	NP_309332	hypothetical protein ECs1305	135.58	3.85E-30	30%	49%	81-430	417-777
					8	cd00079	HELIcC, Helicase superfamily C-terminal domain; associated with DEXDc-, DEAD-, and DEAH-box proteins, yeast initiation factor 4A, Ski2p, and Hepatitis C virus NS3 helicases; this domain is found in a wide variety of helicases and helicase related proteins; may not be an autonomously folding unit, but an integral part of the helicase; 4 helicase superfamilies at present according to the organization of their signature motifs; all helicases share the ability to unwind nucleic acid duplexes with a distinct directional polarity; they utilize the free energy from nucleoside triphosphate hydrolysis to fuel their translocation along DNA, unwinding the duplex in the process	42.22	1.21E-04	23%	40%	302-395	17-123	8	NP_287072	putative helicase	135.58	3.85E-30	30%	49%	81-430	425-785
					9	COG1197	Mif, Transcription-repair coupling factor (superfamily II helicase) [DNA replication, recombination, and repair / Transcription]	42.17	1.34E-04	26%	43%	89-230	607-752	9	YP_00532209	Type III restriction enzyme, res subunit:DEAD/DEAH box helicase, N-terminal	128.26	6.14E-28	29%	46%	51-430	394-790
					10	COG0610	COG0610, Type I site-specific restriction-modification system, R (restriction) subunit and related helicases (Defense mechanisms)	38.12	2.05E-03	22%	36%	95-338	258-517	10	YP_00665630	Type III restriction enzyme, res subunit:DEAD/DEAH box helicase, N-terminal	127.10	1.37E-27	28%	45%	81-430	481-849
N209L	94072-93716	119	12,775	3.90	1	pfam03115	Astro_capsid, Astrovirus capsid protein precursor. This product is encoded by astrovirus ORF2, one of the three astrovirus ORFs (1a, 1b, 2). The 97kD precursor undergoes an intracellular cleavage to form a 79kD protein. Subsequently, extracellular trypsin cleavage yields the three proteins forming the infectious virion.	26.55	7.67E-01	26%	43%	71-117	640-686	1	AAC96525	A157L	93.59	2.12E-18	45%	56%	1-117	1-109
N210L	94496-94128	123	14,411	4.26		No Hit Found							0	No Hit Found	No Hit Found							
N211L	94977-94603	125	13,445	8.66		No Hit Found							0	No Hit Found	No Hit Found							
N214L	95457-95167	97	10,869	9.32		No Hit Found							1	AAC96533	A165L	72.79	3.93E-12	39%	53%	5-95	171-279	
N216L	95974-95516	153	17,584	10.22	1	cd01973	Nitrogenase_VFe_beta_like, Nitrogenase_VFe_delta_like; Nitrogenase VFe protein, beta subunit like. This group contains proteins similar to the beta subunits of the VFe protein of the vanadium-dependent (V-) nitrogenase. Nitrogenase catalyzes the ATP-dependent reduction of dinitrogen (N2) to ammonia. In addition to V-nitrogenase there is a molybdenum (Mo)-dependent nitrogenase and an iron only (Fe-) nitrogenase. The Mo-nitrogenase is the most widespread and best characterized of these systems. These systems consist of component 1 (VFe protein, VFe protein or MoFe protein respectively) and component 2 (Fe protein). MoFe is an alpha2beta2 tetramer, V and Fe-nitrogenases are alpha2beta2delta2 hexamers. The alpha and beta subunits of VFe and FeFe are similar to the alpha and beta subunits of MoFe. For MoFe each alphabeta pair contains one P-cluster (at the alphabeta interface) and, one molecule of iron molybdenum cofactor (FeMoco) contained within the alpha subunit. The Fe protein which has a remarkably identical structure in all these systems. It contains a single Cys	29.52	1.89E-01	41%	49%	47-84	342-383	1	AAC96533	A165L	85.50	5.74E-16	37%	58%	5-118	20-134
N216R	96021-96887	289	33,100	5.82	1	COG5377	COG5377, Phage-related protein, predicted endonuclease [DNA replication, recombination, and repair]	30.01	3.41E-01	23%	32%	30-169	2-148	1	AAC96534	PBCV-1 exonuclease	298.52	1.72E-79	58%	73%	9-253	1-246
N221R	97092-97394	101	10,665	4.11		No Hit Found							2	NP_077549	Esv-1-64	88.58	2.71E-16	27%	50%	21-221	1-191	
N222L	98212-97385	276	30,232	9.99	1	pfam01734	Patatin glycoproteins from plants. The patatin protein accounts for up to 40% of the total soluble protein in potato tubers. Patatin is a storage protein but it also has the enzymatic activity of lipid acyl hydrolase, catalysing the cleavage of fatty acids from membrane lipids. Members of this family have been found also in vertebrates.	116.20	3.75E-27	31%	49%	16-189	1-179	1	AAC96541	similar to E. coli hypothetical protein, corresponds to Swiss-Prot Accession Number P39407	322.40	1.03E-86	57%	77%	5-274	14-288
					2	COG1752	RsaA, Predicted esterase of the alpha-beta hydrolase superfamily [General function prediction only]	102.08	6.31E-23	31%	48%	16-228	14-229	2	YP_00240206	Patatin-like phospholipase family	106.69	8.91E-22	34%	53%	15-185	7-190
					3	COG4667	COG4667, Predicted esterase of the alpha-beta hydrolase superfamily [General function prediction only]	51.42	1.12E-07	24%	40%	16-208	14-202	3	XP_800775	PREDICTED: hypothetical protein XP_795682, partial	99.37	1.42E-19	35%	50%	14-186	76-265
					4	COG3621	COG3621, Patatin [General function prediction only]	31.14	1.52E-01	41%	58%	5-66	6-65	4	YP_133138	hypothetical protein PBPRB1472	90.12	8.63E-17	28%	48%	14-256	6-261
													5	XP_789091	PREDICTED: hypothetical protein XP_783998	89.74	1.13E-16	32%	51%	14-186	86-275	
													6	NP_905966	hypothetical protein PG1879	89.35	1.47E-16	32%	52%	14-186	6-194	
													7	NP_149926	463L	86.66	9.54E-16	29%	54%	14-206	25-215	
													8	AA04389	phospholipase, patatin family	85.89	1.63E-15	30%	50%	15-186	4-194	
													9	AB037620	esterase of the alpha-beta hydrolase superfamily-like	84.73	3.63E-15	30%	50%	15-185	4-193	
													10	YP_142800	patatin-like phospholipase (463L)	83.19	1.06E-14	25%	49%	15-238	57-285	
N223L	99133-98228	302	34,030	4.75	1	cd00180	S_TKc, Serine/Threonine protein kinases, catalytic domain. Phosphotransferases of the serine or threonine-specific kinase subfamily. The enzymatic activity of these protein kinases is controlled by phosphorylation of specific residues in the activation segment of the catalytic domain, sometimes combined with reversible conformational changes in the C-terminal autoinhibitory tail	175.01	8.41E-45	31%	55%	36-295	1-256	1	AAU06282	protein kinase A248R	187.96	3.55E-46	38%	58%	21-295	32-305
					2	smart00220	S_TKc, Serine/Threonine protein kinases, catalytic domain; Phosphotransferases, Serine or threonine-specific kinase subfamily.	175.02	8.98E-45	31%	52%	37-295	1-256	2	AAU06280	protein kinase A248R	187.19	6.05E-46	37%	58%	21-295	32-305
					3	pfam00698	Kinase, Protein kinase domain	164.30	1.48E-41	28%	52%	37-295	1-258	3	AAU06275	protein kinase A248R	187.19	6.05E-46	37%	58%	21-295	11-284
					4	COG0515	SPS1, Serine/threonine protein kinase [General function prediction only / Signal transduction mechanisms / Transcription / DNA replication, recombination, and repair]	117.18	2.27E-27	28%	47%	36-296	1-279	4	AAC96616	PBCV-1 protein kinase	182.19	1.95E-44	37%	58%	24-295	32-304
					5	smart00219	TyrKc, Tyrosine kinase, catalytic domain; Phosphotransferases, Tyrosine-specific kinase subfamily.	91.39	1.42E-19	28%	48%	39-227	3-192	5	AAA87065	serine/threonine protein kinase	180.26	7.39E-44	36%	58%	24-295	28-300
					6	cd00192	TyrKc, Tyrosine kinase, catalytic domain. Phosphotransferases, tyrosine-specific kinase subfamily. Enzymes with TyrKc domains belong to an extensive family of proteins which share a conserved catalytic core common to both serine/threonine and tyrosine protein kinases. Enzymatic activity of tyrosine protein kinases is controlled by phosphorylation of specific tyrosine residues in the activation segment of the catalytic domain or a C-terminal tyrosine (tail) residue with reversible conformational changes.	88.71	7.38E-19	27%	46%	36-227	8-203	6	AAU06274	protein kinase A248R	156.76	8.75E-37	38%	59%	79-295	22-238
					7	COG3642	COG3642, Mch2-dependent serine/threonine protein kinase [Signal transduction mechanisms]	52.93	4.66E-08	28%	43%	112-211	66-180	7	AAU06285	protein kinase A248R	150.60	6.27E-35	37%	58%	81-295	2-217









Table with columns: Gene Name, Genome Position, A.A. length, Peptide Mw, pI, CDD Hit Number, COGs, COG Definition, Bit Score, E-value, % Identity, % Positive, Query from-to, Hit from-to, BLASTp Hit Number, Hit Accession, BLASTp Definition, Bit Score, E-value, % Identity, % Positive, Query from-to, Hit from-to. Contains entries for genes like N288R, N289R, N293R, N299R, N303R, N304R, N307L, N312L, N313L, N315L, N317R, N318L, N320L, N321L, and N323L.



Gene Name	Genome Position	A.A. length	Peptide Mw	pI	CDD Hit Number	COGs	COG Definition	Bit Score	E-value	% Identity	% Positive	Query from-to	Hit from-to	BLASTp Hit Number	Hit Accession	BLASTp Definition	Bit Score	E-value	% Identity	% Positive	Query from-to	Hit from-to		
							Cyt_C5_DNA_methylase. Cytosine-C5 specific DNA methylases; Methyl transfer reactions play an important role in many aspects of biology. Cytosine-specific DNA methylases are found both in prokaryotes and eukaryotes. DNA methylation, or the covalent addition of a methyl group to cytosine within the context of the CpG dinucleotide, has profound effects on the mammalian genome. These effects include transcriptional repression via inhibition of transcription factor binding or the recruitment of methyl-binding proteins and their associated chromatin remodeling factors, X chromosome inactivation, imprinting and the suppression of parasitic DNA sequences. DNA methylation is also essential for proper embryonic development and is an important player in both DNA repair and <i>neurodegeneration</i> .																	
N369L	149048-148017	344	38.430	8.32	1	cd00315		162.40	6.83E-41	30%	43%	5-259	4-271	1	AAC96884	M.CviAII cytosine DNA methyltransferase	447.20	3.96E-124	61%	74%	1-340	1-342		
					2	pfam00145	DNA methylase, C-5 cytosine-specific DNA methylase..	156.24	4.66E-39	29%	44%	5-264	4-267	2	AAC96897	M.CviAIV cytosine DNA methyltransferase	437.96	2.40E-121	61%	76%	1-328	2-332		
					3	COG0270	Dom. Site-specific DNA methylase [DNA replication, recombination, and repair].	108.24	1.21E-24	28%	44%	1-191	3-197	3	AAV84097	CviP1I m5C DNA methyltransferase	309.30	1.29E-82	47%	61%	2-340	16-357		
					4	pfam05478	Prominin. Prominin. The prominins are an emerging family of proteins that among the multipass membrane proteins display a novel topology. Mouse prominin and human prominin (mouse)-like 1 (PROML1) are predicted to contain five membrane spanning domains, with an N-terminal domain exposed to the extracellular space followed by four, alternating small cytoplasmic and large extracellular loops and a cytoplasmic C-terminal domain. The exact function of prominin is unknown although in humans defects in PROM1, the gene coding for prominin cause retinal degeneration.	30.65	3.11E-01	37%	57%	205-240	200-235	4	AAC84006	cytosine methyltransferase	286.19	1.17E-75	44%	58%	2-342	3-358		
					5									5	AAC55063	cytosine methyltransferase	263.08	1.06E-68	40%	56%	5-342	6-362		
					6									6	AAC96987	nonfunctional M.CviAV cytosine DNA methyltransferase	261.92	2.36E-68	40%	56%	5-342	6-362		
					7									7	AAR23218	sp67	101.29	5.34E-20	33%	49%	2-162	6-192		
					8									8	NP_818425	sp127	92.43	2.48E-17	35%	51%	5-155	6-165		
					9									9	NP_150145	putative DNA methylase	92.05	3.24E-17	32%	44%	5-222	4-226		
					10									10	AAC98421	methyl transferase	92.05	3.24E-17	33%	47%	5-170	4-178		
N372L	149874-149227	216	23.956	3.74	1	COG1966	CajA. Carbon starvation protein, predicted membrane protein [Signal transduction mechanism]	28.31	8.02E-01	28%	53%	5-37	2-34	1	AAC96720	Asp/Glu rich; DAEDDDYxxET (2X) negative charge cluster	256.91	3.30E-67	63%	71%	1-214	1-207		
					2									2	YP_142843	unknown	51.22	2.75E-05	32%	58%	135-214	140-221		
N373L	150935-149967	323	36.800	4.01		No Hit Found								1	AAC96725	A357L	164.08	6.10E-39	50%	72%	140-300	86-249		
N377L	151591-150974	206	23.714	5.64		No Hit Found								0	No Hit Found	No Hit Found								
N380R	151657-152352	232	26.010	9.70	1	smart00497	IENR1. Intron encoded nuclease repeat motif. Repeat of unknown function, but possibly DNA-binding via helix-turn-helix motif [Ponting, immunohistochem]	47.43	1.58E-06	37%	54%	175-228	1-53	1	AAC96973	similar to Chlorella virus PBCV-1 ORF A315L, corresponds to GenBank Accession Number U42580	215.31	1.28E-54	50%	63%	1-227	1-224		
					2	smart00465	GIYc, GIY-YIG type nucleases [UR1 domain]; GIY-YIG_Cterm. GIY(X)(Y-1)YIG family of class I homing endonucleases C-terminus (GIY-YIG_Cterm). Homing endonucleases promote the mobility of intron or intein by recognizing and cleaving a homologous allele that lacks the sequence. They catalyze a double-strand break in the DNA near the insertion site of that element to facilitate homing at that site. Class I homing endonucleases are sorted into four families based on the presence of these motifs in their respective N-terminus: LAQLLDKDG, His-Cys box, rH1, and GIY-YIG. This CO contains several but not all members of the GIY-YIG family. The C-terminus of GIY-YIG is a DNA-binding domain which is separated from the N-terminus by a long, flexible linker. The DNA-binding domain consists of a minor-groove binding alpha-helix, and a helix-turn-helix. Some also contain a zinc finger (i.e. I-Tev) which is not required for DNA binding or catalysis, but is a component of the linker and directs the catalytic domain to cleave the homing site at a fixed distance from the intron insertion site.	44.30	1.26E-05	34%	54%	1-91	1-83	2	AA88832	unknown	201.06	2.50E-50	44%	60%	1-228	1-242		
					3	cd00283	GIY-YIG_Cterm. GIY(X)(Y-1)YIG family of class I homing endonucleases C-terminus (GIY-YIG_Cterm). Homing endonucleases promote the mobility of intron or intein by recognizing and cleaving a homologous allele that lacks the sequence. They catalyze a double-strand break in the DNA near the insertion site of that element to facilitate homing at that site. Class I homing endonucleases are sorted into four families based on the presence of these motifs in their respective N-terminus: LAQLLDKDG, His-Cys box, rH1, and GIY-YIG. This CO contains several but not all members of the GIY-YIG family. The C-terminus of GIY-YIG is a DNA-binding domain which is separated from the N-terminus by a long, flexible linker. The DNA-binding domain consists of a minor-groove binding alpha-helix, and a helix-turn-helix. Some also contain a zinc finger (i.e. I-Tev) which is not required for DNA binding or catalysis, but is a component of the linker and directs the catalytic domain to cleave the homing site at a fixed distance from the intron insertion site.	41.14	1.33E-04	46%	61%	111-226	19-113	3	AAC96655	PBCV-1 33kd peptide	172.56	9.51E-42	41%	54%	5-227	7-248		
					4	pfam07453	NUMOD1, NUMOD1 domain..	37.33	1.52E-03	43%	63%	175-205	1-31	4	AAC96862	similar to PBCV-1 ORF A315L, corresponds to GenBank Accession Number M74440	141.35	2.35E-32	41%	55%	1-201	1-196		
					5	pfam01541	GIY-YIG, GIY-YIG catalytic domain. This domain called GIY-YIG is found in the amino terminal region of excinuclease abc subunit c (xrcC), bacteriophage T4 endonucleases segA, segB, segC, segD and segE; it is also found in putative endonucleases encoded by group I introns of fungi and phage. The structure of I-Tev1 a GIY-YIG endonuclease, reveals a novel alpha-beta-fold with a central three-stranded antiparallel beta-sheet flanked by three helices. The most conserved and putative catalytic residues are located on a shallow, concave surface and include a metal coordination site.	36.68	2.71E-03	28%	47%	1-88	1-89	5	YP_293795	putative endonuclease	72.40	1.34E-11	39%	55%	2-108	3-110		
					6	pfam00430	ATP-synth_B. ATP synthase B/B&apos; CF0). Part of the CF0) (base unit) of the ATP synthase. The base unit is thought to translocate protons through membrane (inner membrane in mitochondria, thylakoid membrane in plants, cytoplasmic membrane in bacteria). The B subunits are thought to interact with the stalk of the CF1) subunits. This domain should not be confused with the ab CF1) proteins (in the head of the ATP synthase) which are found in rfam00096.	29.44	3.90E-01	34%	53%	1-31	16-48	6	AAC96906	A539R	53.14	8.41E-06	34%	53%	4-104	34-133		
					7									7	CAA25939	unnamed protein product	50.06	7.12E-05	32%	54%	155-227	75-143		
					8									8	NP_049674	MobB homing endonuclease	50.06	7.12E-05	32%	54%	155-227	112-180		
					9									9	AAC96502	similar to bacteriophage T4 intron-associated endonuclease, corresponds to Swiss-Prot Accession Number P13299	48.91	1.59E-04	34%	52%	2-89	9-95		
					10									10	NP_869393	SegD	48.91	1.59E-04	31%	43%	1-173	1-157		
N383R	152377-155754	1126	123.633	10.87	1	smart00490	HELICc, helicase superfamily c-terminal domain..	37.52	1.02E-02	36%	60%	848-890	38-80	1	AAC96731	similar to chicken vitellogenin II, corresponds to Swiss-Prot Accession Number P02845	835.48	0.00E+00	61%	75%	352-1028	1-651		
					2	pfam00271	Helicase_C. Helicase conserved C-terminal domain. TTis domain family is found in a wide variety of helicases and helicase related proteins. It may be that this is not an autonomously folding unit, but an integral part of the helicase.	36.35	1.96E-02	24%	51%	841-890	27-76	2	AAC96728	A360R	240.74	2.50E-61	55%	68%	4-225	14-239		
					3	cd00079	HELICc. Helicase superfamily c-terminal domain; associated with DEXdc, DEAD-, and DEAH-box proteins, yeast initiation factor 4A, Ski2p, and Hepatitis C virus NS3 helicases; this domain is found in a wide variety of helicases and helicase related proteins; may not be an autonomously folding unit, but an integral part of the helicase; 4 helicase superfamilies are present according to the organization of their signature motifs; all helicases share the ability to unwind nucleic acid duplexes with a distinct directional polarity; they utilize the free energy from nucleoside triphosphate hydrolysis to fuel their translocation along DNA, unwinding the duplex in the reverse.	31.82	4.33E-01	26%	57%	848-890	79-121	3	AAC96729	A361R	130.18	4.76E-28	70%	84%	248-331	2-85		
					4									4	ABF82117	hypothetical protein MIV087L	58.92	1.35E-06	25%	41%	785-938	414-556		
					5									5	YP_142731	putative NTPase I	49.68	8.18E-04	24%	46%	833-941	608-721		
					6									6	YP_142917	helicase conserved C-terminal domain protein	48.21	9.04E-03	30%	51%	832-912	437-519		
					7									7	CAJ57278	putative helicase	46.21	9.04E-03	26%	44%	842-965	318-442		

Gene Name	Genome Position	A.A. length	Peptide Mw	pI	CDD Hit Number	COGs	COG Definition	Bit Score	E-value	% Identity	% Positive	Query from-to	Hit from-to	BLASTp Hit Number	Hit Accession	BLASTp Definition	Bit Score	E-value	% Identity	% Positive	Query from-to	Hit from-to		
N387R	155817-156737	307	35,901	10.26	1	cd00283	GIY-YIG_Cterm, GIYX(10-11)YIG family of class I homing endonucleases C-terminus (GIY-YIG_Cterm). Homing endonucleases promote the mobility of intron or interin by recognizing and cleaving a homologous allele that lacks the sequence. They catalyze a double-strand break in the DNA near the insertion site of that element to facilitate homing at that site. Class I homing endonucleases are sorted into four families based on the presence of these motifs in their respective N-termini: LAGLIDADG, His-Cys box, HN1, and GIY-YIG. The CD contains several but not all members of the GIY-YIG family. The C-terminus of GIY-YIG is a DNA-binding domain which is separated from the N-terminus by a long, flexible linker. The DNA-binding domain consists of a minor-groove binding alpha-helix, and a helix-turn-helix. Some also contain a zinc finger (i.e. I-Tev) which is not required for DNA binding or catalysis, but is a component of the linker and directs the catalytic domain to cleave the homing site at a fixed distance from the intron insertion site.	56.55	4.35E-09	46%	66%	126-187	1-60	1	AA88832	unknown	84.34	5.68E-15	34%	51%	29-198	1-175		
							2	smart00465	GIYc, GIY-YIG type nucleases (URI domain); . GIY-YIG, GIY-YIG catalytic domain. This domain called GIY-YIG is found in the amino terminal region of excinuclease abc subunit c (uvrC), bacteriophage T4 endonucleases segA, segB, segC, segD and segE; it is also found in putative endonucleases encoded by group I introns of fungi and phage. The structure of I-Tev1 a GIY-YIG endonuclease, reveals a novel alpha-beta-fold with a central three-stranded antiparallel beta-sheet flanked by three helices. The most conserved and putative catalytic residues are located on a shallow, concave surface and include a metal coordination site.	31.20	1.69E-01	32%	57%	33-117	6-83	2	AAC96862	similar to PBCV-1 ORF A315L, corresponds to GenBank Accession Number M74440	79.72	1.40E-13	28%	45%	29-276	1-218
							3	pfam01541	GIY-YIG_Cterm, GIYX(10-11)YIG family of class I homing endonucleases C-terminus (GIY-YIG_Cterm). Homing endonucleases promote the mobility of intron or interin by recognizing and cleaving a homologous allele that lacks the sequence. They catalyze a double-strand break in the DNA near the insertion site of that element to facilitate homing at that site. Class I homing endonucleases are sorted into four families based on the presence of these motifs in their respective N-termini: LAGLIDADG, His-Cys box, HN1, and GIY-YIG. The CD contains several but not all members of the GIY-YIG family. The C-terminus of GIY-YIG is a DNA-binding domain which is separated from the N-terminus by a long, flexible linker. The DNA-binding domain consists of a minor-groove binding alpha-helix, and a helix-turn-helix. Some also contain a zinc finger (i.e. I-Tev) which is not required for DNA binding or catalysis, but is a component of the linker and directs the catalytic domain to cleave the homing site at a fixed distance from the intron insertion site.	30.90	1.99E-01	25%	43%	30-112	2-85	3	AAC96973	similar to Chlorella virus PBCV-1 ORF A315L, corresponds to GenBank Accession Number U42580	74.33	5.88E-12	28%	41%	29-276	1-220
							4	AAC96655	PBCV-1 33kd peptide	73.17	1.31E-11	28%	44%	39-276	15-244									
							5	AAC49244	ORF301	66.24	1.60E-09	30%	50%	38-186	83-240									
							6	AAU16837	GIY-YIG catalytic domain containing protein; possible intron encoded endonuclease	63.54	1.04E-08	29%	46%	28-180	2-155									
							7	NP_074951	orf305	57.77	5.70E-07	28%	52%	42-178	85-227									
							8	CAC51107	putative GIY-YIG endonuclease	57.00	9.72E-07	25%	40%	27-274	5-225									
							9	AAC49248	ORF211	55.84	2.16E-06	44%	68%	104-161	86-139									
							10	AAK09365	intron encoded Bmol	53.91	8.23E-06	23%	41%	40-249	14-242									
N389L	157485-156868	206	23,558	10.99	1	smart00465	GIYc, GIY-YIG type nucleases (URI domain); .	34.28	1.14E-02	21%	38%	76-172	1-83	1	AAC96747	A379L	164.85	1.54E-39	44%	60%	12-203	5-203		
							2	COG0322	UvrC, Nuclease subunit of the excinuclease complex [DNA replication, recombination, and repair].	28.33	7.66E-01	41%	47%	72-111	16-50	2	NP_911913	putative probable transcription repressor HOTR	46.60	6.11E-04	25%	40%	42-182	29-193
							3	CAA04677	putative transcription repressor HOTR	45.82	1.04E-03	25%	40%	42-198	26-206									
N391R	157646-158395	250	28,257	8.78	1	cd00283	GIY-YIG_Cterm, GIYX(10-11)YIG family of class I homing endonucleases C-terminus (GIY-YIG_Cterm). Homing endonucleases promote the mobility of intron or interin by recognizing and cleaving a homologous allele that lacks the sequence. They catalyze a double-strand break in the DNA near the insertion site of that element to facilitate homing at that site. Class I homing endonucleases are sorted into four families based on the presence of these motifs in their respective N-termini: LAGLIDADG, His-Cys box, HN1, and GIY-YIG. The CD contains several but not all members of the GIY-YIG family. The C-terminus of GIY-YIG is a DNA-binding domain which is separated from the N-terminus by a long, flexible linker. The DNA-binding domain consists of a minor-groove binding alpha-helix, and a helix-turn-helix. Some also contain a zinc finger (i.e. I-Tev) which is not required for DNA binding or catalysis, but is a component of the linker and directs the catalytic domain to cleave the homing site at a fixed distance from the intron insertion site.	66.95	2.30E-12	46%	63%	101-172	1-72	1	AA88832	unknown	246.90	4.59E-64	51%	70%	1-244	1-240		
							2	smart00497	IENR1, Intron encoded nuclease repeat motif. Repeat of unknown function, but possibly DNA-binding via helix-turn-helix motif (Ponting, unpublished).	42.04	6.10E-05	35%	52%	193-246	1-53	2	AAC96973	similar to Chlorella virus PBCV-1 ORF A315L, corresponds to GenBank Accession Number U42580	213.77	4.31E-54	49%	61%	1-247	1-226
							3	pfam07453	NUMOD1, NUMOD1 domain. GIY-YIG, GIY-YIG catalytic domain. This domain called GIY-YIG is found in the amino terminal region of excinuclease abc subunit c (uvrC), bacteriophage T4 endonucleases segA, segB, segC, segD and segE; it is also found in putative endonucleases encoded by group I introns of fungi and phage. The structure of I-Tev1 a GIY-YIG endonuclease, reveals a novel alpha-beta-fold with a central three-stranded antiparallel beta-sheet flanked by three helices. The most conserved and putative catalytic residues are located on a shallow, concave surface and include a metal coordination site.	37.33	1.74E-03	44%	63%	193-225	1-33	3	AAC96655	PBCV-1 33kd peptide	198.75	1.43E-49	46%	61%	5-244	7-247
							4	pfam01541	GIY-YIG_Cterm, GIYX(10-11)YIG family of class I homing endonucleases C-terminus (GIY-YIG_Cterm). Homing endonucleases promote the mobility of intron or interin by recognizing and cleaving a homologous allele that lacks the sequence. They catalyze a double-strand break in the DNA near the insertion site of that element to facilitate homing at that site. Class I homing endonucleases are sorted into four families based on the presence of these motifs in their respective N-termini: LAGLIDADG, His-Cys box, HN1, and GIY-YIG. The CD contains several but not all members of the GIY-YIG family. The C-terminus of GIY-YIG is a DNA-binding domain which is separated from the N-terminus by a long, flexible linker. The DNA-binding domain consists of a minor-groove binding alpha-helix, and a helix-turn-helix. Some also contain a zinc finger (i.e. I-Tev) which is not required for DNA binding or catalysis, but is a component of the linker and directs the catalytic domain to cleave the homing site at a fixed distance from the intron insertion site.	36.29	3.93E-03	26%	46%	1-88	1-88	4	AAC96862	similar to PBCV-1 ORF A315L, corresponds to GenBank Accession Number M74440	171.79	1.88E-41	45%	62%	1-219	1-196
							5	pfam05118	Asp_Arg_Hydrox, Asparyl/Asparaginyl beta-hydroxylase. Iron (II)-oxoglutarate (2-OG)-dependent oxygenases catalyze oxidative reactions in a range of metabolic processes. Proline 3-hydroxylase hydroxylates proline at position 3, the first of a 2-OG oxygenase catalysing oxidation of a free alpha-amino acid. The structure of proline 3-hydroxylase contains the conserved motifs present in other 2-OG oxygenases including a jelly roll strand core and residues binding iron and 2-oxoglutarate, consistent with divergent evolution within the extended family. This family represent the arginine, asparagine and proline hydroxylases. The asparyl/asparaginyl beta-hydroxylase (EC:1.14.11.16) specifically hydroxylates one aspartic or asparagine residue in certain epidermal growth factor-like domains of a number of proteins.	30.27	2.43E-01	26%	39%	20-91	7-69	5	AAK09365	intron encoded Bmol	82.42	1.50E-14	34%	48%	4-219	5-242
							6	COG4678	COG4678, Muramidase (phage lambda lysozyme) [Carbohydrate transport and metabolism].	28.40	9.87E-01	23%	42%	61-116	24-84	6	NP_899393	SegD	81.65	2.56E-14	33%	46%	1-196	1-205
							7	CAA73995	unnamed protein product	76.64	8.22E-13	31%	49%	14-204	124-311									
							8	AAC49244	ORF301	75.49	1.83E-12	32%	52%	9-159	80-240									
							9	NP_074951	orf305	71.63	2.65E-11	34%	53%	4-162	74-236									
							10	YP_293795	putative endonuclease	67.40	4.99E-10	37%	55%	2-109	3-110									
N395R	158437-159858	474	53,895	6.11	1	pfam04451	Capsid_Hydrov, Iridovirus major capsid protein. This family includes the major capsid protein of iridoviruses, chlorella virus and Spodoptera ascovirus, which are all dsDNA viruses with no RNA stage. This is the most abundant structural protein and can account for up to 45% of virion protein. In Chlorella virus PBCV-1 the major capsid protein is the major protein.	182.43	8.07E-47	29%	45%	81-470	3-422	1	AAC96751	similar to PBCV-1 major capsid protein, corresponds to Swiss-Prot Accession Number P30328	288.50	3.72E-76	52%	72%	148-405	1-257		
							2	pfam03635	Vps35, Vacuolar protein sorting-associated protein 35. Vacuolar protein sorting-associated protein (Vps) 35 is one of around 50 proteins involved in protein trafficking. In particular, Vps35 assembles into a retromer complex with at least four other proteins Vps5, Vps17, Vps20 and Vps29. Vps35 contains a central region of weaker sequence similarity, thought to indicate the presence of at least three domains.	29.51	9.05E-01	21%	39%	86-144	484-546	2	BAE06835	hypothetical major capsid protein	169.09	3.29E-40	31%	49%	81-446	3-401
							3	BAA76601	major capsid protein MCP1	142.12	4.31E-32	30%	45%	81-438	3-380									
							4	BAA22188	major capsid protein Vp54	139.81	2.14E-31	29%	42%	81-469	3-410									
							5	AAC27492	major capsid protein Vp49	139.43	2.79E-31	30%	44%	81-438	3-371									
							6	AAC96798	PBCV-1 major capsid protein Vp54, corresponds to GenBank Accession Number M85052	139.43	2.79E-31	30%	42%	81-469	3-410									
							7	BAA76600	major capsid protein	139.04	3.65E-31	29%	42%	81-469	3-409									
							8	1M4X_C	Chain C, Pbcv-1 Virus Capsid, Quasi-Atomic Model	115.93	3.31E-24	28%	40%	103-469	1-386									
							9	1M3Y_D	Chain D, The Structure Of Major Capsid Protein Of A Large, Lipid Containino, Dna Virus	115.55	4.32E-24	28%	40%	103-469	1-386									
							10	YP_142795	capsid protein	67.40	1.35E-09	31%	46%	284-430	310-465									









Gene Name	Genome Position	A.A. length	Peptide Mw	pl	CDD Hit Number	COGs	COG Definition	Bit Score	E-value	% Identiv	% Positive	Query from-to	Hit from-to	BLASTp Hit Number	Hit Accession	BLASTp Definition	Bit Score	E-value	% Identiv	% Positive	Query from-to	Hit from-to		
							GIY-YIG, GIY-YIG catalytic domain. This domain called GIY-YIG is found in the amino terminal region of excinuclease abc subunit c (xvrc), bacteriophage T4 endonucleases segA, segB, segC, segD and segE; it is also found in putative endonucleases encoded by group I introns of fungi and phage. The structure of I-TevI a GIY-YIG endonuclease, reveals a novel alpha-beta-fold with a central three-stranded antiparallel beta-sheet flanked by three helices. The most conserved and putative catalytic residues are located on a shallow, concave surface and include a metal coordination site.																	
N471L	184168-183284	295	35,080	8.32	1	pfam01541		35.14	1.03E-02	20%	36%	28-112	2-88	1	AAC96862	similar to PBCV-1 ORF A315L, corresponds to GenBank Accession Number M74440	72.02	2.72E-11	30%	49%	30-185	4-166		
					2	smart100465	ABC_SMC4_euk, Eukaryotic SMC4 proteins; SMC proteins are large (approximately 110 to 170 kDa), and each is arranged into five recognizable domains. Amino-acid sequence homology of SMC proteins between species is largely confined to the amino- and carboxy-terminal globular domains. The amino-terminal domain contains a &apostrophes;Walker A&apostrophes; nucleotide-binding domain (GxxGxGKS/T, in the single-letter amino-acid code), which by mutational studies has been shown to be essential in several proteins. The carboxy-terminal domain contains a sequence (the DA-box) that resembles a &apostrophes;Walker B&apostrophes; motif, and a motif with homology to the signature sequence of the ATP-binding cassette (ABC) family of ATPases. The sequence homology within the carboxy-terminal domain is relatively high within the SMC1-SMC4 group, whereas SMC5 and SMC6 show some divergence in both of these sequences. In eukaryotic cells, the proteins are found as heterodimers of SMC1 paired with SMC3, SMC2 with SMC4, and SMC5 with SMC6 (formerly known as Rad18).	33.89	2.28E-02	34%	54%	30-116	4-83	2	AAA88832	unknown	64.31	5.67E-09	29%	45%	30-212	4-180		
					3	cd03274	ABC_SMC4_euk, Eukaryotic SMC4 proteins; SMC proteins are large (approximately 110 to 170 kDa), and each is arranged into five recognizable domains. Amino-acid sequence homology of SMC proteins between species is largely confined to the amino- and carboxy-terminal globular domains. The amino-terminal domain contains a &apostrophes;Walker A&apostrophes; nucleotide-binding domain (GxxGxGKS/T, in the single-letter amino-acid code), which by mutational studies has been shown to be essential in several proteins. The carboxy-terminal domain contains a sequence (the DA-box) that resembles a &apostrophes;Walker B&apostrophes; motif, and a motif with homology to the signature sequence of the ATP-binding cassette (ABC) family of ATPases. The sequence homology within the carboxy-terminal domain is relatively high within the SMC1-SMC4 group, whereas SMC5 and SMC6 show some divergence in both of these sequences. In eukaryotic cells, the proteins are found as heterodimers of SMC1 paired with SMC3, SMC2 with SMC4, and SMC5 with SMC6 (formerly known as Rad18).	32.27	8.15E-02	18%	38%	123-293	885-1060	3	AAC96973	similar to Chlorella virus PBCV-1 ORF A315L, corresponds to GenBank Accession Number U42580	57.38	6.93E-07	39%	51%	30-132	4-105		
					4	pfam04189	elf3_gamma, Eukaryotic initiation factor 3, gamma subunit. elf-3 is a multi-subunit complex that stimulates translation initiation in vitro at several different steps. This family corresponds to the gamma subunit # elf3.	31.52	1.16E-01	23%	47%	150-242	75-165	4	AAC96655	PBCV-1 33kd peptide	56.61	1.18E-06	28%	45%	40-199	19-192		
					5	cd03277	ABC_SMC5_euk, Eukaryotic SMC5 proteins; SMC proteins are large (approximately 110 to 170 kDa), and each is arranged into five recognizable domains. Amino-acid sequence homology of SMC proteins between species is largely confined to the amino- and carboxy-terminal globular domains. The amino-terminal domain contains a &apostrophes;Walker A&apostrophes; nucleotide-binding domain (GxxGxGKS/T, in the single-letter amino-acid code), which by mutational studies has been shown to be essential in several proteins. The carboxy-terminal domain contains a sequence (the DA-box) that resembles a &apostrophes;Walker B&apostrophes; motif, and a motif with homology to the signature sequence of the ATP-binding cassette (ABC) family of ATPases. The sequence homology within the carboxy-terminal domain is relatively high within the SMC1-SMC4 group, whereas SMC5 and SMC6 show some divergence in both of these sequences. In eukaryotic cells, the proteins are found as heterodimers of SMC1 paired with SMC3, SMC2 with SMC4, and SMC5 with SMC6 (formerly known as Rad18).	31.17	1.70E-01	22%	48%	130-240	194-308	5	YP_293795	putative endonuclease	56.61	1.18E-06	39%	55%	38-118	14-95		
							Glycos_transf_2, Glycosyl transferase. Diverse family, transferring sugar from UDP-glucose, UDP-N-acetyl- galactosamine, GDP-mannose or GDP-abequose, to a range of substrates including cellulose, dolichol ribose and fucose acids																	
N472R	184277-186826	850	95,405	6.12	1	pfam00535		49.83	1.32E-06	30%	43%	256-383	2-124	1	AAC96482	A114R	639.80	0.00E+00	64%	77%	373-847	5-477		
					2	COG0463	WcaA, Glycosyltransferases involved in cell wall biogenesis [Cell envelope biogenesis, outer membrane].	47.44	7.61E-06	24%	45%	251-380	2-121	2	AAC96479	A111R	489.96	1.72E-136	58%	75%	1-372	1-379		
					3	COG1215	COG1215, Glycosyltransferases, probably involved in cell wall biogenesis [Cell envelope biogenesis, outer membrane].	44.15	7.72E-05	24%	38%	251-367	53-167	3	NP_872956	possible glycosyltransferase	90.51	3.05E-16	32%	46%	254-426	6-189		
					4	COG1216	COG1216, Predicted glycosyltransferases [General function prediction only].	38.19	4.16E-03	25%	46%	252-369	3-116	4	CAG34747	hypothetical protein	87.81	1.97E-15	29%	45%	1-225	1-225		
					5	pfam03016	Exostosin, Exostosin family. The EXT family is a family of tumour suppressor genes. Mutations of EXT1 on 8q24.1, EXT2 on 11p11-13, and EXT3 on 19p have been associated with the autosomal dominant disorder known as hereditary multiple exostoses (HME). This is the most common known skeletal dysplasia. The chromosomal locations of other EXT genes suggest association with other forms of neoplasia. EXT1 and EXT2 have both been shown to encode a heparan sulphate polymerase with both D-galactosyl (GlcA) and N-acetyl-D-glucosaminoglycan (GlcNAc) transferase activities. The nature of the defect in heparan sulphate biosynthesis in HMF is unclear	35.03	3.45E-02	23%	43%	657-818	177-325	5	ZP_00202013	COG0463: Glycosyltransferases involved in cell wall biogenesis	86.27	5.74E-15	32%	49%	254-426	48-231		
N477R	186874-187335	154	16,607	8.18		No Hit Found																		
N480R	187361-187714	118	12,745	11.93		No Hit Found																		
N482R	187960-188919	320	36,440	8.67	1	smart100507	HNhc, HNH nucleases; .	29.67	5.70E-01	36%	45%	127-168	10-52	1	AAC96845	Lys-, Arg-rich; contains eukaryotic putative RNA-binding region RNP-1 signature; similar to PBCV-1 ORF A267L, corresponds to GenBank Accession Number U42580	188.73	2.27E-46	35%	47%	9-312	3-305		
					2	cd00085	HNhc, HNH nucleases; HNH endonuclease signature which is found in viral, prokaryotic, and eukaryotic proteins. The alignment includes members of the large group of homing endonucleases, yeast intron 1 protein, MutS, as well as bacterial colicins, ovocins, and anaretoxins..	28.89	8.81E-01	30%	43%	127-172	11-57	2	AAC96857	Lys-, Glu-rich	177.56	5.24E-43	34%	47%	9-312	3-305		
					3	COG1110	COG1110, Reverse gyrase [DNA replication, recombination, and repair].	28.74	9.54E-01	30%	52%	4-27	692-715	3	YP_142777	unknown	73.17	1.96E-11	38%	56%	194-295	251-349		
					4																			
N484L	189482-189174	103	10,874	10.74	1	pfam05854	MC1, Non-histone chromosomal protein MC1. This family consists of archaeal chromosomal protein MC1 sequences which protect DNA against thermal denaturation..	42.35	1.23E-05	37%	53%	5-68	2-70	1	AAC96805	similar to Methanotrix chromosomal protein MC1A, corresponds to Swiss-Prot Accession Number P15251	134.81	8.23E-31	69%	81%	5-99	6-100		
					2																			
					3																			
					4																			
					5																			
					6																			
					7																			
					8																			
N485R	189557-189775	73	7,645	10.67		No Hit Found																		



Gene Name	Genome Position	A.A. length	Peptide Mw	pI	CDD Hit Number	COGs	COG Definition	Bit Score	E-value	% Identity	% Positive	Query from-to	Hit from-to	BLASTp Hit Number	Hit Accession	BLASTp Definition	Bit Score	E-value	% Identity	% Positive	Query from-to	Hit from-to
N531L	207897-209900	666	73,134	8.43		No Hit Found								1	AAC98403	contains Pro-rich Px motif, PAFK (19X); similar to Arabidopsis anter-specific Pro-rich protein, corresponds to Swiss-Prot Accession Number P40602	50.06	3.42E-04	28%	45%	174-291	30-134
N533L	208755-207985	257	29,166	8.44		No Hit Found								2	BAA11344	DNA binding protein	51.22	1.54E-04	26%	36%	133-289	95-261
														1	AAC96377	A9R	222.25	1.28E-56	63%	79%	85-250	8-173
														2	AAC96972	similar to Chlorella virus PBCV-1 ORF A450R, corresponds to GenBank Accession Number U42580	129.03	1.48E-28	31%	51%	1-251	4-255
														3	AAC96818	similar to PBCV-1 ORF A275R, encoded by GenBank Accession Number U42580	118.24	2.61E-25	29%	50%	5-249	2-246
														4	AAC96643	similar to PBCV-1 ORF A79R, corresponds to GenBank Accession Number U17055	113.24	8.39E-24	28%	49%	5-248	5-248
														5	AAC96545	A177R	91.66	2.61E-17	25%	44%	5-252	5-245
														6	AAC96447	A79R	91.28	3.41E-17	27%	45%	5-228	2-218
														7	AAC96971	similar to Chlorella virus PBCV-1 ORF A450R, corresponds to GenBank Accession Number U42580	85.89	1.43E-15	64%	90%	5-55	2-52
														8	AAU06304	hypothetical protein A275R	58.92	1.88E-07	27%	46%	85-248	4-167
														9	AAU06301	hypothetical protein A275R	57.77	4.18E-07	26%	46%	85-248	4-167
10	AAU06302	hypothetical protein A275R	57.00	7.13E-07	28%	47%	117-251	1-134														
N535L	213044-209160	1295	130,191	6.38	1	COG0196	RibF, FAD synthase [Coenzyme metabolism].	35.26	5.65E-02	25%	39%	1026-1143	164-274	1	AAC96907	A540L	101.29	2.76E-19	24%	36%	7-562	523-1052
														2	ABE69167	A540L	50.45	5.59E-04	26%	41%	9-170	389-544
														3	YP_589431	hypothetical protein Acid345_0352	52.37	1.47E-04	23%	46%	28-198	438-599
														4	AA048404	unknown protein	137.53	3.45E-30	37%	52%	1080-1292	50-280
														5	ZP_01257209	haemalectinin family protein	80.49	5.04E-13	23%	35%	30-465	24-412
														6	ZP_00533161	Hep Haq	60.46	5.40E-07	22%	45%	12-209	54-249
														7	YP_214529	possible T4-like proximal tail fiber	85.11	2.05E-14	21%	35%	12-652	362-960
														8	ZP_0094573	Collagen alpha (Z1) chain precursor	72.40	1.37E-10	25%	33%	335-730	736-1083
														9	ZP_00532602	Hep Haq	49.68	9.53E-04	24%	38%	40-209	684-867
														10	ZP_00950302	outer membrane protein	51.22	3.28E-04	25%	45%	28-178	640-784
														0	No Hit Found	No Hit Found						
N540L	214063-213113	317	36,864	10.55	1	COG0436	COG0436, Aspartate/tyrosine/aromatic aminotransferase [Amino acid transport and metabolism].	29.12	7.54E-01	28%	44%	33-87	124-174	0	No Hit Found	No Hit Found						
N542L	215171-214032	380	41,731	10.39	1	COG3007	COG3007, Uncharacterized paraquat-inducible protein B [Function unknown].	29.53	7.19E-01	37%	47%	282-348	74-133	1	AAC96985	STKPP (11x); similar to Gossypium Pro-rich wall protein, corresponds to GenBank Accession Number U04267	181.80	3.60E-44	46%	60%	159-378	163-384
N546L	215578-215141	146	16,059	7.00	1	cd01286	Deoxycytidylate deaminase. Deoxycytidylate deaminase domain. Deoxycytidylate deaminase catalyzes the deamination of dCMP to dUMP, providing the nucleotide substrate for thymidylate synthase. The enzyme binds Zn <sup>2+</sup> , which is required for catalytic activity. The activity of the enzyme is allosterically regulated by the ratio of dCTP to dTTP not only in eukaryotic cells but also in T-even phage-infected Escherichia coli with dCTP acting as an activator and dTTP as an inhibitor	128.08	3.69E-31	39%	55%	9-135	4-131	1	AAC96936	similar to Vibrio fischeri dCMP deaminase, corresponds to Swiss-Prot Accession Number P33668	201.45	7.16E-51	65%	81%	4-144	2-142
														2	COG2131	ComEB, Deoxycytidylate deaminase [Nucleotide transport and metabolism].	104.27	5.66E-24	36%	54%	9-141	12-148
														3	pfam00383	dCMP_cyt_deam, Cytidine and deoxycytidylate deaminase zinc-binding region..	86.57	9.86E-19	41%	60%	8-113	5-100
														4	cd00786	cytidine deaminase-like. Cytidine and deoxycytidylate deaminase zinc-binding region. The family contains cytidine deaminases, nucleoside deaminases, deoxycytidylate deaminases and riboflavin deaminases. Also included are the apoBec family of mRNA editing enzymes. All members are Zn dependent. The zinc ion in the active site plays a central role in the proposed catalytic mechanism, activating a water molecule to form a hydroxide ion that performs a nucleophilic attack on the substrate.	58.87	2.71E-10	29%	50%	14-113	6-92
														5	cd01285	nucleoside deaminase. Nucleoside deaminases include adenosine, guanine and cytosine deaminases. These enzymes are Zn dependent and catalyze the deamination of nucleosides. The zinc ion in the active site plays a central role in the proposed catalytic mechanism, activating a water molecule to form a hydroxide ion that performs a nucleophilic attack on the substrate. The functional enzyme is a homodimer. Cytosine deaminase catalyzes the deamination of cytosine to uracil and ammonia and is a member of the pyrimidine salvage pathway. Cytosine deaminase is found in bacteria and fungi but is not present in mammals; for this reason, the enzyme is currently of interest for antimicrobial drug design and gene therapy applications against tumors. Some members of this family are RNA-specific adenosine deaminases that generate inosine at the first position of their anticodon (position 34) of specific tRNAs; this modification is thought to enlarge the codon recognition capacity during protein synthesis. Other members of the family are guanine deaminases which deaminate inosine in vivo as part of the utilization of inosine	56.43	1.38E-09	32%	55%	10-120	1-100
														6	cd01284	Riboflavin deaminase reductase. Riboflavin-specific deaminase. Riboflavin biosynthesis protein RibD (Diaminohydroxyphosphoribosylaminopyrimidine deaminase) catalyzes the deamination of 2,5-diamino-6-ribosylamino-4(3H)-pyrimidinone 5&apos;-phosphate, which is an intermediate step in the biosynthesis of riboflavin. The ribG gene of Bacillus subtilis and the ribD gene of E. coli are bifunctional and contain this deaminase domain and a reductase domain which catalyzes the subsequent reduction of the ribosyl side chain	55.25	2.72E-09	38%	56%	28-113	21-93
														7	COG0590	CumB, Cytosine/adenosine deaminases [Nucleotide transport and metabolism / Transition, ribosomal structure and biogenesis].	54.57	5.04E-09	34%	50%	1-113	3-104
														8	COG0117	RibD, Pyrimidine deaminase [Coenzyme metabolism].	54.10	7.39E-09	28%	51%	1-113	1-100
9	YP_293777	putative deoxycytidylate deaminase	97.44	1.46E-19	50%	67%	21-113	35-134														
N548R	215695-216513	273	30,481	8.08	1	cd00180	S_TKc, Serine/Threonine protein kinases, catalytic domain. Phosphotransferases of the serine or threonine-specific kinase subfamily. The enzymatic activity of these protein kinases is controlled by phosphorylation of specific residues in the activation segment of the catalytic domain, sometimes combined with reversible conformational changes in the C-terminal adenine/uracil tail	150.74	1.47E-37	29%	50%	15-265	1-256	1	AAC96657	similar to PBCV-1 serine/threonine protein kinase, corresponds to GenBank Accession Number U14660	136.73	7.88E-31	33%	49%	4-265	6-279
														2	smart00220	S_TKc, Serine/Threonine protein kinases, catalytic domain. Phosphotransferases. Serine or threonine-specific kinase subfamily	144.97	8.97E-36	28%	48%	16-265	1-256
														3	pfam00069	Kinase, Protein kinase domain.	143.89	1.70E-35	30%	49%	16-265	1-258
														4	COG0515	SPT, Serine/threonine protein kinase [General function prediction only / Signal transduction mechanisms / Transcription / DNA replication, recombination and repair]	106.78	2.54E-24	26%	44%	16-269	2-282
														5	smart00219	TyrKc, Tyrosine kinase, catalytic domain; Phosphotransferases. Tyrosine-specific kinase subfamily.	74.44	1.48E-14	24%	44%	17-258	2-250
														6	cd00192	TyrKc, Tyrosine kinase, catalytic domain. Phosphotransferases; tyrosine-specific kinase subfamily. Enzymes with TyrKc domains belong to an extensive family of proteins which share a conserved catalytic core common to both serine/threonine and tyrosine protein kinases. Enzymatic activity of tyrosine protein kinases is controlled by phosphorylation of specific tyrosine residues in the activation segment of the catalytic domain or a C-terminal tyrosine (tail) residue with reversible conformational changes	68.68	7.32E-13	24%	43%	13-261	6-264
														7	COG3894	COG3894, Uncharacterized metal-binding protein [General function prediction only].	35.69	7.13E-03	21%	35%	130-247	105-227
														8	pfam01636	APH, Phosphotransferase enzyme family. This family consists of bacterial antibiotic resistance proteins, which confer resistance to various aminoglycosides; they include 3&apos;-aminoglycoside 3&apos;-phosphotransferase or kanamycin kinase / neomycin-kanamycin phosphotransferase and streptomycin 3&apos;-phosphotransferase or streptomycin 3&apos;-phosphotransferase. The aminoglycoside phosphotransferases inactivate aminoglycoside antibiotics via phosphorylation. This family also includes homoserine kinase. This family is related to fructosamine kinase rfam1881	31.66	1.10E-01	36%	52%	116-157	156-198
														9	COG2334	COG2334, Putative homoserine kinase type II (protein kinase fold) [General function prediction only]	30.70	1.92E-01	27%	43%	114-158	183-227
														9	AA161666	AI2g30360/T9D9.17	91.66	2.91E-17	28%	47%	22-264	27-276

Gene Name	Genome Position	A.A. length	Peptide Mw	pI	CDD Hit Number	COGs	COG Definition	Bit Score	E-value	% Identiv	% Positive	Query from-to	Hit from-to	BLASTp Hit Number	Hit Accession	BLASTp Definition	Bit Score	E-value	% Identiv	% Positive	Query from-to	Hit from-to
					10	COG3173	COG3173, Predicted aminoglycoside phosphotransferase [General function, non-membrane]	29.61	3.88E-01	42%	52%	125-157	195-226	10	AAU06282	protein kinase A248R	91.28	3.79E-17	25%	45%	14-268	46-308
N549R	216631-219810	1060	117,600	9.53	1	smart00433	TOP2c, TopoisomeraseII; Eukaryotic DNA topoisomerase II, GyrB, ParE	453.09	7.69E-128	32%	49%	50-610	1-589	1	AAU95770	topoisomerase II	1889.39	0.00E+00	90%	92%	1-1058	1-1058
					2	smart00434	TOP4c, DNA Topoisomerase IV; Bacterial DNA topoisomerase IV, GyrA, ParC	392.63	1.11E-109	33%	50%	626-1017	1-423	2	AAC96932	PBCV-1 DNA topoisomerase II	1338.55	0.00E+00	63%	75%	3-1056	2-1061
					3	cd00187	TOP4c, DNA Topoisomerase, subtype IIA; domain A&apoc; bacterial DNA topoisomerase IV (C subunit, ParC), bacterial DNA gyrase (A subunit, GyrA), mammalian DNA topoisomerase II. DNA topoisomerase are essential enzymes that regulate the conformational changes in DNA topology by catalysing the concerted breakage and rejoining of DNA strands during normal cellular growth.	362.96	8.47E-101	33%	51%	644-1017	1-401	3	NP_584718	DNA TOPOISOMERASE II	877.09	0.00E+00	47%	64%	5-1017	8-1030
					4	COG0187	GyrB, Type IIA topoisomerase (DNA gyrase/topo II, topoisomerase IV), B subunit (DNA nicking, recombination, and repair)	319.39	1.17E-87	27%	43%	2-626	8-635	4	NP_189031	TOPII (TOPOISOMERASE II); ATP binding / DNA binding / DNA	846.27	0.00E+00	46%	63%	5-1000	35-1072
					5	pfam00521	DNA topoisolV, DNA gyrase/topoisomerase IV, subunit A.	271.27	3.49E-73	30%	45%	646-1017	1-397	5	AAW40881	DNA topoisomerase II, putative	844.34	0.00E+00	45%	61%	5-1000	112-1153
					6	COG0188	GyrA, Type IIA topoisomerase (DNA gyrase/topo II, topoisomerase IV), A subunit (DNA replication, recombination, and repair)	251.33	3.96E-67	23%	44%	617-1015	2-428	6	XP_467311	putative DNA topoisomerase II	834.71	0.00E+00	45%	60%	5-1000	32-1097
					7	cd03365	TOPRIM, TopoIIA, TopoIIA, Topoisomerase-primase (TOPRIM) nucleotidyl transferase/hydrolase domain of the type found in proteins of the type IIA family of DNA topoisomerases similar to Saccharomyces cerevisiae Topoisomerase II. TopoIIA enzymes cut both strands of the duplex DNA to remove (relax) both positive and negative supercoils in DNA. These enzymes covalently attach to the 5&apoc; ends of the cut DNA, separate the free ends of the cleaved strands, pass another region of the duplex through this gap, then rejoin the ends. These proteins also catenate/ decatenate duplex rings. The TOPRIM domain has two conserved motifs, one of which centers at a conserved glutamate and the other one at two conserved aspartates (DxD). This glutamate and two aspartates, cluster together to form a highly acid surface patch. The conserved glutamate may act as a general base in strand joining and as a general acid in strand cleavage by topoisomerases. The DxD motif may co-ordinate Mg2+, a cofactor required for full catalytic function.	142.67	2.18E-34	66%	78%	403-521	1-120	7	BAE06274	topoisomerase II	833.56	0.00E+00	45%	60%	5-1000	35-1078
					8	cd01030	TOPRIM, TopoIIA, TopoIIA, Topoisomerase-primase (TOPRIM) nucleotidyl transferase/hydrolase domain of the type found in proteins of the type IIA family of DNA topoisomerases similar to Saccharomyces cerevisiae Topoisomerase II. TopoIIA enzymes cut both strands of the duplex DNA to remove (relax) both positive and negative supercoils in DNA. These enzymes covalently attach to the 5&apoc; ends of the cut DNA, separate the free ends of the cleaved strands, pass another region of the duplex through this gap, then rejoin the ends. These proteins also catenate/ decatenate duplex rings. The TOPRIM domain has two conserved motifs, one of which centers at a conserved glutamate and the other one at two conserved aspartates (DxD). The conserved glutamate may act as a general base in strand joining and as a general acid in strand cleavage by topoisomerases. The DxD motif may co-ordinate Mg2+, a cofactor required for full catalytic function.	123.67	1.00E-28	53%	63%	403-521	1-115	8	BAD86854	DNA topoisomerase II	833.17	0.00E+00	43%	60%	5-1000	103-1163
					9	cd03366	TOPRIM, TopoIIA, GyrB, TOPRIM, TopoIIA, GyrB, topoisomerase-primase (TOPRIM) nucleotidyl transferase/hydrolase domain of the type found in proteins of the type IIA family of DNA topoisomerases similar to the Escherichia coli GyrB subunit. TopoIIA enzymes cut both strands of the duplex DNA to remove (relax) both positive and negative supercoils in DNA. These enzymes covalently attach to the 5&apoc; ends of the cut DNA, separate the free ends of the cleaved strands, pass another region of the duplex through this gap, then rejoin the ends. These proteins also catenate/ decatenate duplex rings. DNA gyrase is more effective at relaxing supercoils than decatenating DNA. DNA gyrase in addition inserts negative supercoils in the presence of ATP. The TOPRIM domain has two conserved motifs, one of which centers at a conserved glutamate and the other one at two conserved aspartates (DxD). The conserved glutamate may act as a general base in strand joining and as a general acid in strand cleavage by topoisomerases. The DxD motif may co-ordinate Mg2+, a cofactor required for full catalytic function.	82.10	2.96E-16	44%	57%	403-510	1-105	9	AAN85208	DNA topoisomerase II	831.63	0.00E+00	45%	62%	5-1000	27-1065
					10	pfam00204	DNA_gyraseB, DNA gyrase B. This family represents the second domain of DNA gyrase B which has a ribosomal S5 domain 2-like fold. This family is structurally related to PF01119.	62.57	2.23E-10	27%	44%	237-382	14-168	10	XP_751245	DNA topoisomerase II	828.17	0.00E+00	44%	61%	2-996	55-1094
N569R	219871-220755	295	33,581	10.47	1	cd00283	GIY-YIG_Cterm, GIYX(10-11)YIG family of class I homing endonucleases C-termus (GIY-YIG_Cterm). Homing endonucleases promote the mobility of intron or intron by recognizing and cleaving a homologous allele that lacks the sequence. They catalyze a double-strand break in the DNA near the insertion site of that element to facilitate homing at that site. Class I homing endonucleases are sorted into four families based on the presence of these motifs in their respective N-termini: LAGLIDADG, His-Cys box, HNH, and GIY-YIG. This CD contains several but not all members of the GIY-YIG family. The C-terminus of GIY-YIG is a DNA-binding domain which is separated from the N-terminus by a long, flexible linker. The DNA-binding domain consists of a minor-groove binding alpha-helix, and a helix-turn-helix. Some also contain a zinc finger (i.e. I-TevI) which is not required for DNA binding or catalysis, but is a component of the linker and directs the catalytic domain to cleave the homing site at a fixed distance from the intron insertion site.	55.39	9.05E-09	41%	61%	181-275	1-100	1	AAA88832	unknown	194.90	2.78E-48	40%	55%	1-289	1-240
					2	smart00497	ENR1, Intron encoded nuclease repeat motif. Repeat of unknown function, but possibly DNA-binding via helix-turn-helix motif (Ponting, unpublished).	48.98	7.50E-07	42%	60%	239-291	1-53	2	AAC96973	similar to Chlorella virus PBCV-1 ORF A315L, corresponds to GenBank Accession Number U42590	190.66	5.25E-47	39%	53%	1-290	1-224
					3	pfam07453	NUMOD1, NUMOD1 domain.	44.65	1.50E-05	48%	67%	239-272	1-34	3	AAC96655	PBCV-1 33kd peptide	173.33	8.67E-42	35%	52%	3-290	6-248
					4	smart00465	GIYc, GIY-YIG type nucleases (URI domain).	40.06	3.56E-04	32%	54%	1-72	1-70	4	AAC96862	similar to PBCV-1 ORF A315L, corresponds to GenBank Accession Number M74440	48.14	4.21E-04	42%	67%	179-234	97-151
					5	COG1693	COG1693, Uncharacterized protein conserved in archaea [Function unknown]	35.64	7.69E-03	29%	48%	108-186	13-90	5	NP_899393	SegD	85.11	3.10E-15	29%	46%	1-241	1-197
					6	pfam01995	DUF128, Domain of unknown function DUF128. This archaebacterial protein family has no known function. The domain is found duplicated in rna number	35.33	1.03E-02	27%	48%	108-186	14-91	6	YP_253795	putative endonuclease	72.02	2.72E-11	44%	58%	2-84	3-85
					7	pfam01541	GIY-YIG, GIY-YIG catalytic domain. This domain called GIY-YIG is found in the amino terminal region of excinuclease abo subunit c (uvrC), bacteriophage T4 endonucleases segA, segB, segC, segD and segE; it is also found in putative endonucleases encoded by group I introns of fungi and phage. The structure of I-TevI a GIY-YIG endonuclease, reveals a novel alpha-beta-fold with a central three-stranded antiparallel beta-sheet flanked by three helices. The most conserved and putative catalytic residues are located on a shallow, concave surface and include a metal coordination site.	33.98	2.43E-02	33%	56%	1-59	1-64	7	AAC49244	ORF301	71.63	3.55E-11	28%	48%	4-170	76-236
					8	COG3069	DcuC, C4-dicarboxylate transporter [Energy production and conversion].	31.79	1.05E-01	26%	43%	23-65	187-229	8	NP_074951	orf305	70.86	6.06E-11	32%	49%	4-204	74-262
					9	pfam06342	DUF1057, Protein of unknown function (DUF1057). This family consists of several Caenorhabditis elegans specific proteins of unknown function.	31.43	1.41E-01	31%	48%	118-202	200-284	9	AAC49248	ORF211	55.84	2.02E-06	28%	47%	129-290	49-208
					10									10	AAT53588	group I intron GIY-YIG endonuclease	49.68	1.45E-04	25%	43%	4-205	5-197
N561R	220777-221190	138	15,751	11.06	1	cd02698	Pepitidase_C1A_CathepsinX, Cathepsin X, the only papain-like lysosomal cysteine peptidase exhibiting carboxymonopeptidase activity. It can also act as a carboxypeptidase, like cathepsin B, but has been shown to preferentially cleave substrates through a monopeptidyl carboxypeptidase pathway. The propeptide region of cathepsin X, the shortest among papain-like peptidases, is covalently attached to the active site cysteine in the inactive form of the enzyme. Little is known about the biological function of cathepsin X. Some studies point to a role in early tumorigenesis. A more recent study indicates that cathepsin X expression is restricted to immune cells suggesting a role in phagocytosis and the regulation of the immune response.	26.82	9.48E-01	68%	73%	9-35	40-62	1	AAC96929	A577L	108.61	6.33E-23	55%	62%	26-130	2-104











Gene Name	Genome Position	A.A. length	Peptide Mw	pI	CDD Hit Number	COGs	COG Definition	Bit Score	E-value	% Identity	% Positive	Query from-to	Hit from-to	BLASTp Hit Number	Hit Accession	BLASTp Definition	Bit Score	E-value	% Identity	% Positive	Query from-to	Hit from-to																																																													
N670L	257268-256465	268	29.615	6.63	1	COG0571	Rnc, dsRNA-specific ribonuclease [Transcription].	172.76	3.27E-44	35%	51%	38-263	4-234	4	NP_078699	Thiol oxidoreductase	62.00	6.88E-09	33%	55%	2-99	21-115																																																													
							5	XP_503294	hypothetical protein	62.00	6.88E-09	33%	50%	9-111	88-186																																																																				
							6	XP_451806	unnamed protein product	60.08	2.61E-08	31%	50%	14-111	85-178																																																																				
							7	CAA48192	ERV1	58.54	7.60E-08	31%	48%	14-110	19-111																																																																				
							8	NP_011543	Flavin-linked sulphydryl oxidase localized to the mitochondrial intermembrane space, has a role in the maturation of cytosolic iron-sulfur proteins; ortholog of human hepatocysteinyl (ALR); Erv1o	58.54	7.60E-08	31%	48%	14-110	91-183																																																																				
							9	AAL98767	ORF043L	56.61	2.89E-07	34%	52%	7-99	5-96																																																																				
							10	AAX82354	thiol oxidoreductase	56.23	3.77E-07	34%	52%	7-99	5-96																																																																				
							2	AAC96831	similar to Bacillus ribonuclease III, corresponds to Swiss-Prot Accession Number P51833	133.65	6.43E-30	35%	53%	33-258	9-241																																																																				
							2	YP_445467	ribonuclease III	133.65	6.43E-30	35%	53%	33-258	9-241																																																																				
							3	ZP_00590199	Ribonuclease III	128.26	2.70E-28	36%	54%	45-255	47-265																																																																				
4	ZP_00532592	Ribonuclease III	127.49	4.61E-28	33%	53%	16-255	21-259																																																																											
5	ABR23018	Ribonuclease III	122.87	1.14E-26	35%	55%	58-255	47-252																																																																											
6	ZP_00591208	Ribonuclease III	122.48	1.48E-26	34%	52%	45-256	44-266																																																																											
7	ZP_00528534	Ribonuclease III	122.48	1.48E-26	33%	54%	41-255	58-281																																																																											
8	ZP_00511103	Ribonuclease III	121.32	3.30E-26	34%	54%	45-257	42-264																																																																											
9	ZP_00661649	Ribonuclease III	121.32	3.30E-26	32%	50%	35-257	19-254																																																																											
10	AAM73335	ribonuclease III	119.78	9.62E-26	32%	52%	30-262	15-260																																																																											
N672L	257494-257279	72	7.981	9.71	1	COG2433	COG2433, Uncharacterized conserved protein [Function unknown].	28.34	2.34E-01	18%	55%	15-59	421-465	0	No Hit Found	No Hit Found																																																																			
N674R	257683-259627	655	75.058	5.23	1	COG3378	COG3378, Predicted ATPase [General function prediction only].	70.01	7.92E-13	24%	37%	281-593	146-445	1	AAC96824	contains ATP/GTP-binding site motif A	871.69	0.00E+00	65%	81%	20-648	22-650																																																													
							2	pfam03288	PoxVirus D5 protein-like. This family includes D5 from Poxviruses which is necessary for viral DNA replication, and is a nucleic acid independent nucleoside triphosphatase. Members of this family are also found outside of poxviruses..	63.73	6.04E-11	22%	39%	290-614	3-313	2	NP_077594	EsV-1-109	278.49	5.80E-73	35%	54%	190-607	141-565																																																											
N679R	259658-260536	293	31.095	5.47	No Hit Found	No Hit Found																																																																													
																							N682L	261298-260618	227	26.441	7.60	1	pfam01108	Tissue fac, Tissue factor..	28.90	5.70E-01	25%	46%	144-223	89-176	1	AAC96923	A568L	71.25	2.88E-11	26%	53%	41-194	29-173																																						
																														N684L	261677-261282	132	14.749	7.24	No Hit Found	No Hit Found																																															
																																																				N685R	261742-262269	176	19.819	7.15	1	COG5098	COG5098, Chromosome condensation complex Condensin, subunit D2 [Chromatin structure and dynamics / Cell division and chromosome maintenance]	30.44	1.17E-01	21%	44%	12-135	333-444	1	AAC96926	A572R	243.82	1.75E-63	65%	85%	4-163	11-170									
																																																											N687L	263042-262260	261	28.789	4.23	1	cd00577	PCNA, Proliferating Cell Nuclear Antigen (PCNA) domain found in eukaryotes and archaea. These polymerase processivity factors play a role in DNA replication and repair. PCNA encircles duplex DNA in its central cavity, providing a DNA-bound platform for the attachment of the polymerase. The trimeric PCNA ring is structurally similar to the dimeric ring formed by the DNA polymerase processivity factors in bacteria (beta subunit DNA polymerase III holoenzyme) and in bacteriophages (catalytic subunits in T4 and RB69). This structural correspondence further substantiates the mechanistic connection between eukaryotic and prokaryotic DNA replication that has been suggested on biochemical grounds. PCNA is also involved with proteins involved in cell cycle processes such as DNA repair and apoptosis. Many of these proteins contain a highly conserved motif known as the PIP-box [PCNA interacting protein box] which contains the sequence Qxx[LM]p[DFY].	142.72	3.31E-35	27%	51%	21-251	5-247	1	AAC96927	similar to Periwinkle PCNA, corresponds to GenBank Accession Number X55052	231.49	2.18E-59	44%	67%	4-259	4-264		
																																																																		2	pfam02747	PCNA_C, Proliferating cell nuclear antigen, C-terminal domain, N-terminal and C-terminal domains of PCNA are topologically identical. Three PCNA molecules are tightly associated to form a closed ring encircling duplex DNA.	65.73	4.87E-12	28%	49%	135-250	3-128	2	CAE67843	Hypothetical protein CBG13430	125.56	1.68E-27	29%	50%	15-257	2-260
																																																																		3	COG0592	DnaN, DNA polymerase sliding clamp subunit (PCNA homolog) [DNA recombination, and repair].	59.93	2.92E-10	18%	37%	1-251	47-323	3	EAR2594	proliferating cell nuclear antigen (pcna)	117.86	3.51E-25	28%	48%	15-255	2-259
																																																																		4	pfam00705	PCNA_N, Proliferating cell nuclear antigen, N-terminal domain, N-terminal and C-terminal domains of PCNA are topologically identical. Three PCNA molecules are tightly associated to form a closed ring encircling duplex DNA.	49.88	3.27E-07	25%	55%	25-112	12-100	4	CAA38636	proliferating cell nuclear antigen	117.09	5.99E-25	30%	49%	20-257	7-260
																																																																		5	pfam04139	FadG, FadG, FadG is required for transient cell-cycle arrests and transcriptional induction of DNA repair in response to DNA damage	34.29	1.67E-02	22%	47%	15-73	3-61	5	AAC48257	Pcna (proliferating cell nuclear antigen) homolog protein 1	117.09	5.99E-25	30%	51%	48-256	1-226
																																																																		6	COG1355	COG1355, Predicted dioxygenase [General function prediction only].	30.25	2.93E-01	17%	44%	21-153	110-227	6	XP_502661	hypothetical protein	115.16	2.26E-24	25%	52%	25-258	12-260
7	XP_743255	proliferating cell nuclear antigen (PCNA)	112.85	1.13E-23	29%	51%	21-251	7-261																																																																											
8	CAA38893	proliferating cell nuclear antigen	112.46	1.48E-23	30%	49%	25-258	12-262																																																																											
10	AAG24908	proliferating cell nuclear antigen	112.08	1.93E-23	30%	49%	25-257	12-261																																																																											
N689L	263595-263089	169	19.586	10.14	1	COG1052	LdhA, Lactate dehydrogenase and related dehydrogenases [Energy production and conversion / Coenzyme metabolism / General function prediction only].	29.44	2.57E-01	22%	41%	97-148	93-147	1	AAC96928	A575L	112.46	5.42E-24	35%	66%	31-166	30-167																																																													
N690R	263679-264704	342	37.532	8.60	1	smart00494	ChitBD2, Chitin-binding domain type 2 ;	38.96	8.41E-04	40%	50%	278-321	9-49	1	AAC96701	PLPRNLLL (4X), SPSPSKP (3X)	343.58	6.11E-93	70%	80%	7-220	1-213																																																													
							2	pfam03067	Chitin_bind_3, Chitin binding domain. This domain is found associated with a wide variety of cellulose binding domain. This domain however is a chitin binding domain. This domain is found in isolation in baculoviral spherulins and spinolins, protein of unknown function.	38.43	1.20E-03	22%	29%	18-216	1-211	2	AAC96700	a332L	88.20	4.63E-16	74%	90%	280-322	1-43																																																											
							3	pfam01607	CBM_14, Chitin binding Peritrophin-A domain. This domain is called the Peritrophin-A domain and is found in chitin binding proteins particularly peritrophic matrix proteins of insects and animal chitinases. Copies of the domain are also found in some bacteriophages. Relevant references that describe proteins with this domain include. It is an extracellular domain that contains six conserved cysteines that probably form three disulphide bridges. Chitin binding has been demonstrated for a protein containing only two of these domains	36.57	4.88E-03	41%	62%	290-321	18-47	3	EAA01148	ENSANGP0000018413	74.71	5.30E-12	27%	41%	25-217	4-202																																																											
							4	COG3397	COG3397, Uncharacterized protein conserved in bacteria [Function unknown].	35.41	1.19E-02	22%	33%	11-222	8-213	4	XP_966436	PREDICTED: similar to CG4367-PA isoform 1	72.79	2.01E-11	33%	41%	15-217	4-209																																																											



Gene Name	Genome Position	A.A. length	Peptide Mw	pI	CDD Hit Number	COGs	COG Definition	Bit Score	E-value	% Identiv	% Positive	Query from-to	Hit from-to	BLASTp Hit Number	Hit Accession	BLASTp Definition	Bit Score	E-value	% Identiv	% Positive	Query from-to	Hit from-to
N720L	275415-273727	563	63.379	11.37	1	COG0515	SPS1, Serine/threonine protein kinase [General function prediction only / Signal transduction mechanisms / Transcription / DNA replication, recombination, and repair].	43.61	7.23E-05	13%	28%	84-456	11-365	1	AAC96947	RPOT-like (Rx)	460.69	6.86E-128	44%	62%	2-561	6-577
					2	cd0180	S_TKc_Serine/Threonine protein kinases, catalytic domain. Phosphotransferases of the serine or threonine-specific kinase subfamily. The enzymatic activity of these protein kinases is controlled by phosphorylation of specific residues in the activation segment of the catalytic domain, sometimes combined with reversible conformational changes in the C-terminal autoregulatory tail.	42.89	1.04E-04	30%	46%	53-238	6-151	2	AAC96646	similar to bovine cyclin I, corresponds to Swiss-Prot Accession Number P35662	291.20	7.20E-77	39%	60%	1-401	9-424
					3	pfam00069	Pkinase, Protein kinase domain..	41.04	3.63E-04	28%	45%	53-236	5-148	3	AAC96650	similar to PBCV-1 ORF A34R, corresponds to GenBank Accession Number U17055	285.03	5.16E-75	36%	56%	1-453	23-465
					4	pfam01636	APH1, Phosphotransferase enzyme family. This family consists of bacterial antibiotic resistance proteins, which confer resistance to various aminoglycosides they include- aminoglycoside 3&apos;-phosphotransferase or kanamycin kinase / neomycin-kanamycin phosphotransferase and streptomycin 3&apos;-phosphotransferase or streptomycin 3&apos;-phosphotransferase. The aminoglycoside phosphotransferases inactivate aminoglycoside antibiotics via phosphorylation. This family also includes homoserine kinase. This family is related to fructosyltransferase pfam00001.	40.90	4.29E-04	41%	74%	207-233	171-198	4	XP_944436	PREDICTED: similar to CG10953-PA	62.39	5.44E-08	33%	55%	349-446	76-173
					5	COG2334	COG2334, Putative homoserine kinase type II (protein kinase fold) [General function prediction only]	39.94	7.47E-04	33%	52%	207-240	200-233	5	XP_932523	PREDICTED: similar to serine/arginine repetitive matrix 2	55.84	5.09E-06	35%	52%	375-456	97-178
					6	smart00220	S_TKc_Serine/Threonine protein kinases, catalytic domain. Phosphotransferases. Serine or threonine-specific kinase subfamily.	39.81	1.02E-03	25%	48%	53-238	5-150	6	XP_729588	dentin phosphoryn	47.75	1.39E-03	36%	51%	348-429	315-396
					7	COG0661	AarF, Predicted unusual protein kinase [General function prediction only].	38.39	2.48E-03	31%	59%	204-254	284-333	7	AAK54495	neurofilament triplet H1-like protein	57.00	2.29E-06	36%	50%	327-456	93-221
					8	smart00090	RIO, RIO-like kinase..	38.29	2.62E-03	28%	45%	166-250	122-211	8	XP_742899	hypothetical protein PC108912.00.0	47.05	1.30E-03	28%	55%	362-431	42-111
					9	pfam01163	RIO1, RIO1 family. This family of proteins are related to eukaryotic type protein kinases..	37.50	3.98E-03	28%	50%	162-252	78-170	9	XP_427855	PREDICTED: similar to p87, partial	45.44	6.88E-03	29%	50%	375-446	86-157
					10	cd00142	PI3K family. Phosphoinositide 3-kinase, catalytic domain; Phosphoinositide 3-kinase isoforms participate in a variety of processes, including cell motility, the Ras pathway, vesicle trafficking and secretion, and apoptosis. These homologues may be either lipid kinases and/or protein kinases; the former phosphorylate the 3-position in the inositol ring of inositol phospholipids. The ataxia telangiectasia-mutated gene product, the targets of rapamycin (TOR) and the DNA-dependent kinase have not been found to possess lipid kinase activity. Some of this family possesses PI-4 kinase activities.	37.08	6.09E-03	38%	49%	209-252	147-192	10	AAC96402	similar to E. coli LPS core biosynthesis protein, corresponds to Swiss-Prot Accession Number P27240	50.06	2.79E-04	20%	38%	26-313	29-307
N724R	275496-276452	319	37.146	10.16		No Hit Found								1	AAC96948	similar to Variola virus orf E10L, corresponds to Swiss-Prot Accession Number P33901	284.65	3.03E-75	44%	65%	1-310	1-314
														2	YP_142754	S/T protein kinase, similar to Paramecium bursaria chlorella virus 1 A617R	58.15	4.61E-07	29%	51%	130-246	230-358
N725L	276823-276443	127	13.956	3.77		No Hit Found								1	AAC96949	A618L	75.87	4.60E-13	54%	79%	56-123	65-125
N727L	277600-276950	217	24.102	4.26	1	cd00829	SCP-x thiolase. Thiolase domain associated with sterol carrier protein (SCP)-x isoform and related proteins. SCP-2 has multiple roles in intracellular lipid circulation and metabolism. The N-terminal presequence in the SCP-x isoform represents a peroxisomal 3-ketacyl-Coa thiolase specific for branched-chain acyl CoAs, which is proteolytically cleaved from the sterol carrier protein.	31.44	8.80E-02	31%	52%	118-172	7-61	1	AAC96950	A619L	51.60	2.13E-05	24%	38%	1-215	1-237
N731L	277945-277649	99	11.140	8.64		No Hit Found								1	AAC96951	similar to Synecocystis orf 90, corresponds to GenBank Accession Number D69002	64.31	1.39E-09	34%	50%	1-95	1-81
														2	AAC96952	A635R	50.45	2.07E-05	30%	51%	1-91	1-82
N732L	278380-278027	118	12.691	8.65		No Hit Found								1	AAC96952	A621L	130.18	2.06E-29	52%	71%	4-116	5-117
N733R	278451-281159	903	99.656	6.17	1	COG0498	Uup, ATPase components of ABC transporters with duplicated ATPase domains [General function prediction only].	280.22	6.30E-76	27%	45%	300-848	2-530	1	AAC96981	Chlorella virus CVK2 translation elongation factor-3 homolog, refer to GenBank Accession Number D16505	1106.66	0.00E+00	65%	77%	37-901	55-918
					2	cd03221	ABC_EF-3, ABCF_EF-3 Elongation factor 3 (EF-3) is a cytosolic protein required by fungal ribosomes for in vitro protein synthesis and for in vivo growth. EF-3 stimulates the binding of the EF-1-GTP-aa-tRNA methyl complex to the ribosomal A site by facilitated release of the deacylated tRNA from the E site. The reaction requires ATP hydrolysis. EF-3 contains two ATP binding sequence (NBS) motifs. NBSI is sufficient for the intrinsic ATPase activity. NBSII is essential for the ribosome-stimulated function.	174.24	5.16E-44	39%	56%	302-492	1-191	2	A48779	translation elongation factor EF-3 homolog - Chlorella virus CVK2	1011.52	0.00E+00	61%	74%	37-885	272-1120
					3	pfam00005	ABC_tran, ABC transporter. ABC transporters for a large family of proteins responsible for translocation of a variety of compounds across biological membranes. ABC transporters are the largest family of proteins in many completely sequenced bacteria. ABC transporters are composed of two copies of this domain and two copies of a transmembrane domain pfam00664. These four domains may belong to a single polypeptide, or belong in different polypeptide chains.	120.40	8.69E-28	30%	48%	327-491	1-182	3	XP_445123	unnamed protein product	757.67	0.00E+00	46%	62%	35-900	137-1043
					4	cd00267	ABC_ATPase, ABC (ATP-binding cassette) transporter nucleotide-binding domain; ABC transporters are a large family of proteins involved in the transport of a wide variety of different compounds, like sugars, ions, peptides and more complex organic molecules. The nucleotide binding domain shows the highest similarity between all members of the family. ABC transporters are a subset of nucleotide hydrolases that contain a signature motif, Q-loop, and H-loop/switch region in addition to the Walker A motif/loop and Walker B motif commonly found in a number of ATP- and GTP-binding and hydrolyzing proteins..	114.78	3.59E-26	29%	47%	306-491	4-207	4	BA433959	translation elongation factor3	756.13	0.00E+00	46%	62%	35-900	137-1043
					5	cd03230	ABC_DR_subfamily_A, This family of ATP-binding proteins belongs to a multisubunit transporter involved in drug resistance (BcrA and DnrA), modulation, lipid transport, and lambicid immunity. In bacteria and archaea, these transporters usually include an ATP-binding protein and one or two integral membrane proteins. Eukaryote systems of the ABCA subfamily display ABC domains that are quite similar to this family. The ATP-binding domain shows the highest similarity between all members of the ABC transporter family. ABC transporters are a subset of nucleotide hydrolases that contain a signature motif, Q-loop, and H-loop/switch region in addition to the Walker A motif/loop and Walker B motif commonly found in a number of ATP- and GTP-binding and hydrolyzing proteins.	112.09	2.44E-25	29%	47%	311-488	10-205	5	CAA78282	translation elongation factor 3	753.82	0.00E+00	45%	62%	29-901	136-1050
					6	COG1131	CmaA, ABC-type multidrug transport system, ATPase component [Defense mechanisms].	103.13	1.29E-22	28%	48%	312-487	16-208	6	XP_711404	translation elongation factor 3	753.44	0.00E+00	45%	62%	29-901	136-1050
					7	cd03225	ABC_cobalt_transport_domain1, Domain I of the ATPase component of a cobalt transport family found in both bacteria and archaea. This ABC transporter subfamily is involved cobalt transport as part of the cobalamin biosynthetic pathway. Cobalamin is derived from uroporphyrinogen III, a precursor of heme, siroheme and chlorophylls, and a cobalt ion is chelated in the center of the molecules. The genes necessary for cobalamin production are organized into a single operon in S. typhimurium. In addition to genes known to encode enzymes catalyzing steps of the cobalamin biosynthetic pathway, the products of cbiQ, cbiO and cbiN, were proposed to constitute a cobalt uptake system since CbiN and CbiQ are integral membrane proteins and CbiO is an ABC ATPase. However, direct evidence supporting this idea is lacking..	102.50	1.87E-22	28%	46%	306-491	4-209	7	CAA22654	SPCC417.08	751.90	0.00E+00	45%	63%	35-898	142-1046
					8	COG1121	Znuc, ABC-type Mn/Zn transport systems, ATPase component [Inorganic ion transport and metabolism].	101.06	5.09E-22	29%	50%	306-493	9-216	8	XP_711356	translation elongation factor 3	751.13	0.00E+00	45%	62%	29-901	136-1050









Gene Name	Genome Position	A.A. length	Peptide Mw	pI	CDD Hit Number	COGs	COG Definition	Bit Score	E-value	% Identity	% Positive	Query from-to	Hit from-to	BLASTp Hit Number	Hit Accession	BLASTp Definition	Bit Score	E-value	% Identity	% Positive	Query from-to	Hit from-to
N847R	319064-319798	245	27,064	8.51		No Hit Found																
														1	AAC96734 A366L	similar to Chlorella virus PBCV-1 ORF A450R, corresponds to GenBank Accession Number U42580	155.61	1.35E-36	38%	59%	6-237	14-251
														2	AAC96972	similar to Chlorella virus PBCV-1 ORF A450R, corresponds to GenBank Accession Number U42580	61.62	2.66E-08	32%	46%	2-151	8-144
														3	AAC96818	similar to PBCV-1 ORF A275R, encoded by GenBank Accession Number U42580	56.61	8.55E-07	37%	51%	1-72	1-81
														4	AAC96643	similar to PBCV-1 ORF A79R, corresponds to GenBank Accession Number U17055	55.45	1.91E-06	25%	40%	1-237	4-199
														5	AAC96447 A70R	similar to Chlorella virus PBCV-1 ORF A450R, corresponds to GenBank Accession Number U42580	55.07	2.49E-06	27%	40%	1-237	1-189
														6	AAC96971	similar to Chlorella virus PBCV-1 ORF A450R, corresponds to GenBank Accession Number U42580	54.30	4.25E-06	48%	59%	1-47	1-47
														7	AAC96545 A177R		51.22	3.59E-05	42%	63%	1-47	4-50