#### Bioinformatics CS300

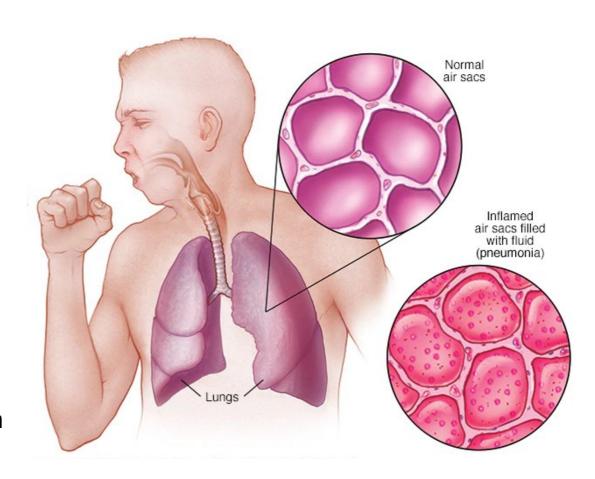
Blast, Substitution Matrices and Protein Alignments (Chap 4 and 5 in textbook)

Fall 2019
Oliver BONHAM-CARTER



#### Pneumonia

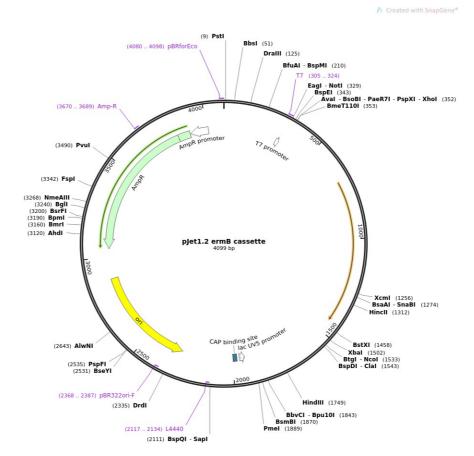
- Pneumonia is an infection that inflames the air sacs in one or both lungs. The air sacs may fill with fluid or pus (purulent material), causing cough with phlegm or pus, fever, chills, and difficulty breathing. A variety of organisms, including bacteria, viruses and fungi, can cause pneumonia.
- A classic sign of bacterial pneumonia is a cough that produces thick, blood-tinged or yellowish-greenish sputum with pus.





#### Human Pathogen Inquiry: The *ermB* gene

 An erythromycin-resistance gene from Streptococcus agalactiae, a gram-positive bacterial species commonly associated with the udders of cows, causing mastitis (i.e., inflammation of breast tissue that sometimes involves an infection and may cause fever)

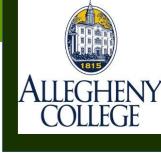




#### Pneumonia and ermB

- Drug resistant: Erythromycin is a macrolide antibiotic used to treat bacterial infections
- Resistance is due to the *ermB* gene which has been noted in the bacteria, *Streptococcus* pneumonia – a common cause of bacterial pneumonia.

#### Horizontal Gene Transfer?



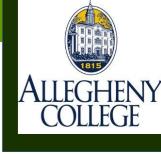
- This type of pneumonia is not believed to have always been resistant to drugs.
- Could the resistance gene have come from another bacteria via HGT?
- How could we check what other bacterial organisms have a specific allele for the gene that effectively resists drugs?
- We will use Blast for this task.

BLAST

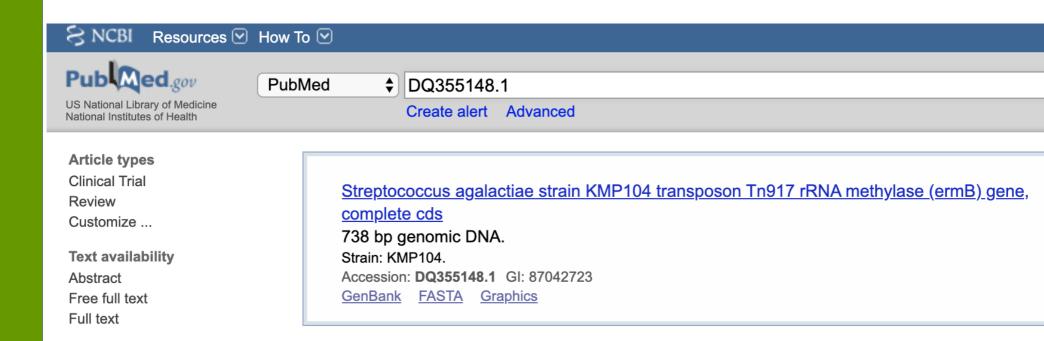
BLAST

BLAST



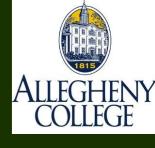


- Locate the Accession number, DQ355148.1, on http://www.pubmed.gov
- Streptococcus agalactiae strain KMP104 transposon Tn917 rRNA methylase (ermB) gene, complete cds



https://www.ncbi.nlm.nih.gov/nuccore/87042723/





Send to: -

GenBank

Streptococcus agalactiae strain KMP104 transposon Tn917 rRNA methylase (ermB) gene, complete cds

GenBank: DQ355148.1

FASTA Graphics

#### Go to: 🔽

LOCUS DQ355148 738 bp DNA linear BCT 13-FEB-2006

DEFINITION Streptococcus agalactiae strain KMP104 transposon Tn917 rRNA

methylase (ermB) gene, complete cds.

ACCESSION DQ355148 VERSION DO355148.1

KEYWORDS

SOURCE Streptococcus agalactiae
ORGANISM Streptococcus agalactiae

Bacteria; Firmicutes; Bacilli; Lactobacillales; Streptococcaceae;

Streptococcus.

REFERENCE 1 (bases 1 to 738)

AUTHORS Puopolo, K.M., Klinzing, D.C., Lin, M.P., Yesucevitz, D.L. and

Cieslewicz, M.J.

TITLE A Composite Transposon Responsible for ErmB-Mediated Erythromycin

Resistance in Group B Streptococcus

JOURNAL Unpublished

REFERENCE 2 (bases 1 to 738)

AUTHORS Puopolo, K.M., Klinzing, D.C., Lin, M.P., Yesucevitz, D.L. and

Cieslewicz, M.J.

TITLE Direct Submission

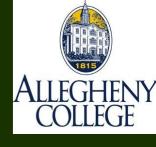
JOURNAL Submitted (06-JAN-2006) Channing Laboratory, Brigham and Women's

Hospital, 181 Longwood Avenue, Boston, MA 02115, USA

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Recent activity

Q DQ355148.1

Streptococcus

transposon Tr

GenBank -

#### Streptococcus agalactiae strain KMP104 transposon Tn917 rRNA (ermB) gene, complete cds

GenBank: DQ355148.1

FASTA Graphics

#### Go to: ✓

LOCUS DQ355148 738 bp DNA linear BCT 13-FEB-2006

DEFINITION Streptococcus agalactiae strain KMP104 transposon Tn917 rRNA

methylase (ermB) gene, complete cds.

ACCESSION DQ355148 VERSION DQ355148.1

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Resistance in Group B Streptococcus

JOURNAL Unpublished

REFERENCE 2 (bases 1 to 738)

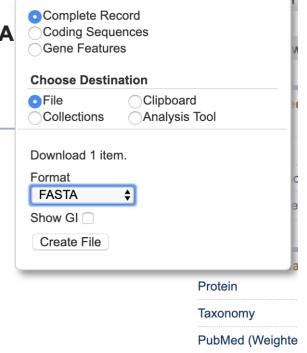
AUTHORS Puopolo, K.M., Klinzing, D.C., Lin, M.P., Yesucevitz, D.L. and

Cieslewicz, M.J.

TITLE Direct Submission

JOURNAL Submitted (06-JAN-2006) Channing Laboratory, Brigham and Women's

Hospital, 181 Longwood Avenue, Boston, MA 02115, USA

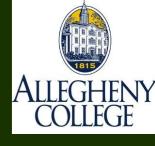


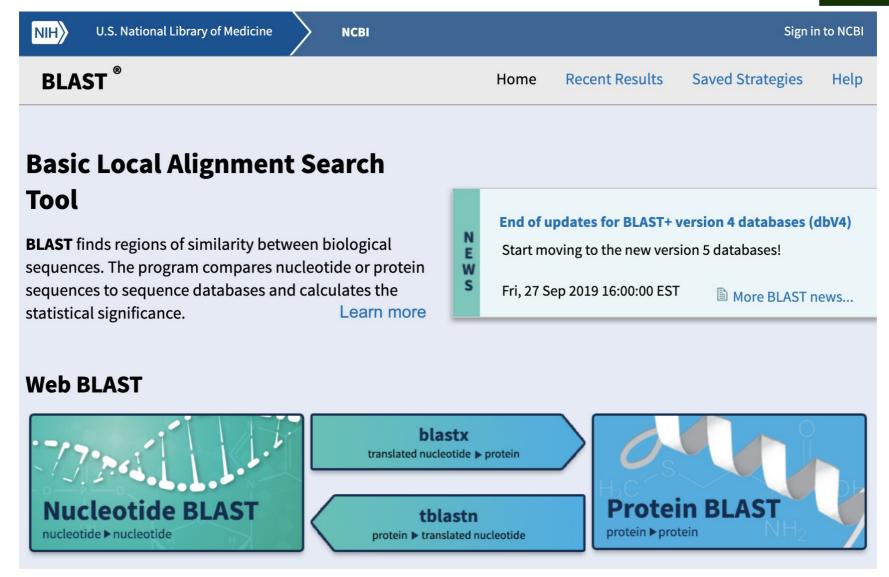
Send to: -





#### **Blast Website**

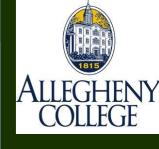




https://blast.ncbi.nlm.nih.gov/Blast.cgi

#### Run The Query





blastn	<u>blastp</u>	<u>blastx</u>	<u>tblastn</u>	tblastx						
			В	LASTN pr	ograms search n	nucleotide dat	abases using a	nucleotide	query. more	
Ent	ter Que	ery Sequ	uence							
Enter	access	ion num	ber(s), g	i(s), or F	ASTA sequenc	ce(s) 😡		<u>Clear</u>		
			ococcus	agalactia	e strain KMP10	04 transpos	on Tn917 rRN	A methyla	se (ermB) gene,	
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					TATTGAATTAG					
					ATTCACCAAGA					
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Cho	oose S	earch S	Set							
Datab	oase		Human	genomi	c + transcript	Mouse ger	nomic + transc	ript O0th	er (Tr J.c.):	
			Nucleotic	de collecti	on (nr/nt)			<b>○ ©</b>		

Use database: *Nucleotide* collection (nr/nt)

### Results

1363 1363 100%

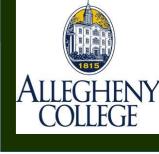
0.0 100.00% MH626525.1



	criptions	Graphic Summary	Alignments	Taxonomy							
Seq	uences p	roducing significant a	lignments	Download ~	Mana	ige Co	lumns	× (	Show 1	00 🕶	
<b>2</b> 9	select all 1	.00 sequences selected			<u>Gen</u>	<u>Bank</u>	<u>Graph</u>	<u>iics</u>	<u>Distance t</u>	ree of results	
		Des	scription		Max Score	Total Score	Query Cover	E value	Per. Ident	Accession	
	Staphylococ	ccus aureus strain VGC1 chromo	osome, complete genom	<u>e</u>	1363	1363	100%	0.0	100.00%	CP039448.1	
	Enterococcu	us durans strain VREdu plasmid	pSULI, complete seque	nce	1363	1363	100%	0.0	100.00%	CP043327.1	
~	Enterococcu	us durans strain VREdu chromos	<u>some</u>		1363	1363	100%	0.0	100.00%	CP042597.1	
<b>~</b>	Enterococcu	us faecalis EnGen0107 strain B5	<u>i94 plasmid p2, complete</u>	e sequence	1363	1363	100%	0.0	100.00%	CP041740.1	
<b>~</b>	Enterococcu	us faecalis strain 4928STDY707	1263 genome assembly,	chromosome: 1	1363	1363	100%	0.0	100.00%	LR607346.1	
	Enterococcu	us faecium strain N56454 plasmi	id unnamed, complete s	<u>equence</u>	1363	1363	100%	0.0	100.00%	CP040905.1	
	Enterococcu	us avium strain 352 plasmid unn	amed, complete sequen	<u>ce</u>	1363	1363	100%	0.0	100.00%	CP034168.1	
	Listeria mon	ocytogenes hypothetical protein	<u>, IS1216 transposase, 3</u>	-aminoglycoside o-phosp	1363	1363	100%	0.0	100.00%	MK490828.1	
<b>~</b>	Enterococcu	us faecium isolate E8407 genom	e assembly, plasmid: 2		1363	1363	100%	0.0	100.00%	LR536659.1	
<b>~</b>	Enterococcu	us faecium SMVRE20 plasmid p	SMVRE20S DNA, comp	lete genome	1363	1363	100%	0.0	100.00%	AP019410.1	
<b>✓</b>	Enterococcu	ıs faecium strain 37BA plasmid ր	oEf37BA, complete sequ	<u>ience</u>	1363	1363	100%	0.0	100.00%	MG957432.1	
	Enterococcu	us faecium strain FSIS1608820 p	olasmid pFSIS1608820,	complete sequence	1363	2668	100%	0.0	100.00%	CP028728.1	
	Streptococc	us pneumoniae isolate GPS_HK	<u>_21-sc-2296565 genom</u>	ne assembly, chromosome	1363	1363	100%	0.0	100.00%	LR216058.1	

Synthetic construct clone pEP1237, complete sequence

#### Scores



#### Max Score

- The score of the best matching segment for local alignment, not global

#### Total Score

 The total scores of all matching segments found (same as max score if there is only one matching segment)

#### Query Coverage

- The percentage of the query sequence that aligned to some part of the match.

#### E-Value

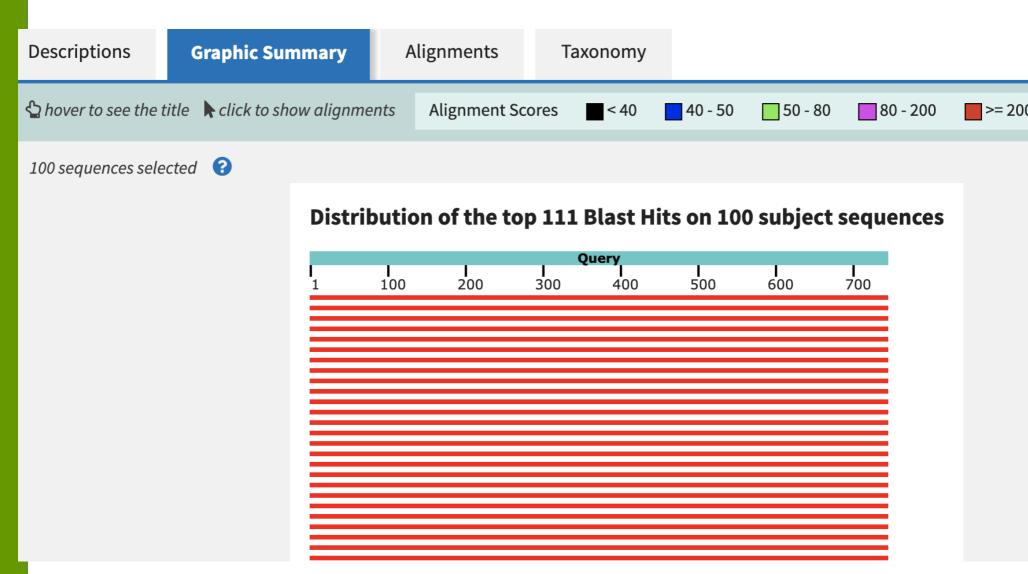
 A statistical measure evaluating how likely it is that a match this good could occur by chance. Lower e-scores indicate that both sequences are truly similar and are not similar by chance alone. Identical sequences have e-scores of zero.

#### Max Indent

 The percentage of nucleotides that are identical between the query and the target sequences within the matching regions.

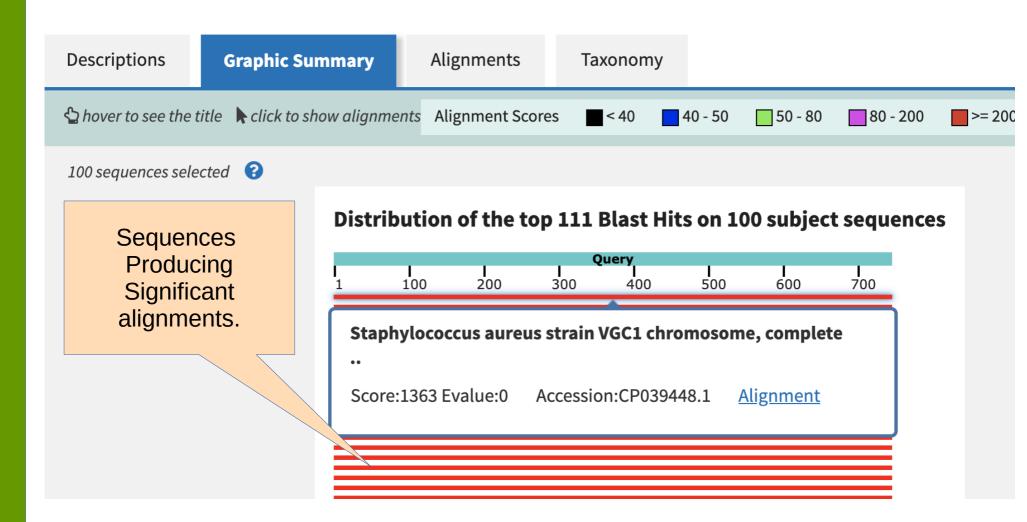
#### Results













#### Results

#### **Streptococcus suis strain SC216 ICESsuSC216 sequence**

Sequence ID: MK359991.1 Length: 54396 Number of Matches: 2

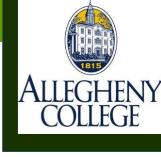
	Range 1	l: <b>15</b> 9	998	to 16451 GenB	ank <u>Graphics</u>		▼ Next Match
	Score			Expect	Identities	Gaps	Strand
	839 bits	s(454	)	0.0	454/454(100%)	0/454(0%)	Plus/Plus
	Query	1	1	AACAGGTAACG	TGAATTAGACA	GTCATCTATTCAACTTA	TCGTCAGAAAA
An Id	entica	J	00	TAACO	GTCTATTGAATTAGACA	GTCATCTATTCAACTTA	TCGTCAGAAAA
•	ence ir ther's		58		FCGTGTCACTTTAATTC 		
ger	nome			AACAGAGGTAT	 	TTCCTTACCATTTAAGC	ACACAAATTATI
	Sbjct	161	18	AACAGAGGTAT	raaaartgttgggaata	TTCCTTACCATTTAAGC	ACACAAATTATI
	Query	181		AAGTGGTTTTT	rgaaagccgtgcgtctg 	ACATCTATCTGATTGTT 	'GAAGAAGGATT( 
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	Sbjct	162	38	AGCGTACCTTC	ĠĠĀŢĀŢŢĊĀĊĊĠĀĀĊĀĊ	ŢŖĠĠĠŢŢĠĊŢĊŢŢĠĊŖĊ	zÁCTCÁÁGTCTCC
	Query	301		AGCAATTGCTT	FAAGCTGCCAGCGGAAT(	GCTTTCATCCTAAACCA 	AAAGTAAACAG1
	Sbjct	162	98	AGCAATTGCT	raagctgccagcgaat	ĠĊŦŦŦĊĂŦĊĊŦĂĂĂĊĊĀ	AAAGTAAACAGT



#### Back to HGT?

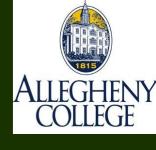
- Typically, researchers allow for a 95% similarity between genes found between *unrelated* organisms.
- Here, we may conclude that HGT is a good hypothesis but more research must be done to determine whether there was a chance for two organisms to be close enough to each other to share genetic material.











- Investigate a gene of resistance: ermA
- Questions:
  - What does this gene do? (hint: see Genbank record)
  - About how many other organisms appear to have traces of the same gene sequence?
  - What is the closest match? Which organism? What *e*-score?



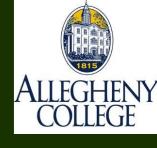


GitHub Activity Repository: https://classroom.github.com/a/DuBJW7yi

Due at 12:15 on 17 Oct. 2019

Make a directory: **act1**Workfile: **act1/blastWork.md** 

#### Predicted Protein Results?



Results may not have been experimentally observed, DNA can be translated to produce this protein.

The reading frame of the DNA might produce a different protein than this one

♣ Download ➤ GenPept Graphics

PREDICTED: serine/threonine-protein kinase PINK1, mitochondria

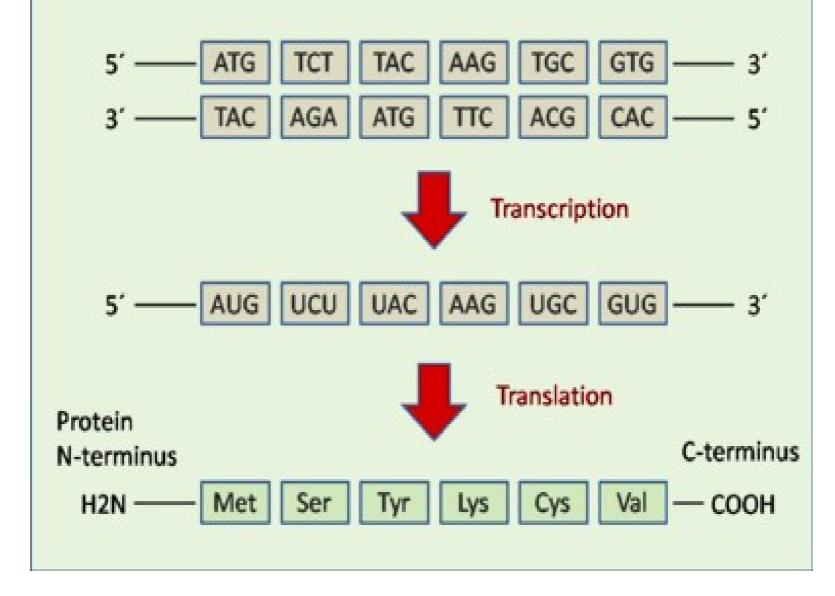
Sequence ID: XP\_014893419.1 Length: 575 Number of Matches: 1

Range 1: 1 to 334 GenPept Graphics

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Sbjct	241	QITLD	GRFGVL	LPRRVS	AHPNV	IRVYRA	AFTA	DVP	LLPG	AQE	EY	2DV	/LE
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#### The central dogma of molecular biology

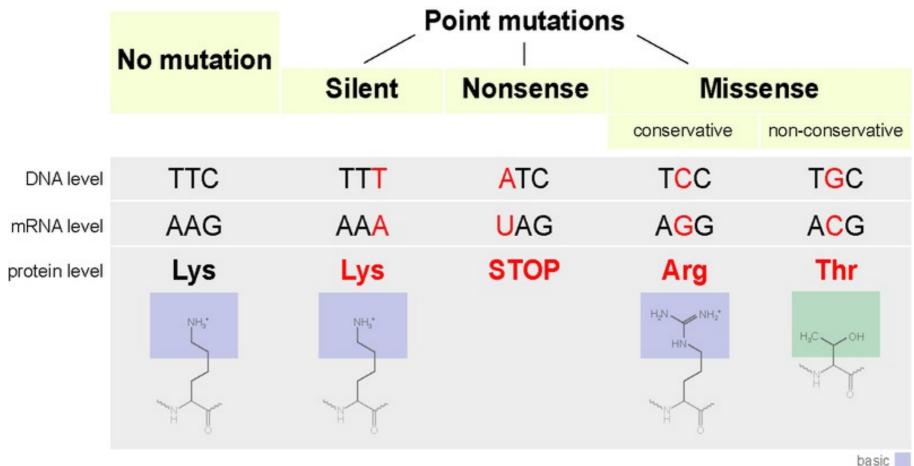




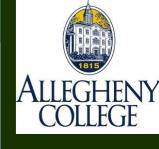
polar

#### More About Silent Mutations

 Redundant codons mean ~1/3 of DNA mutations often do not alter protein sequence

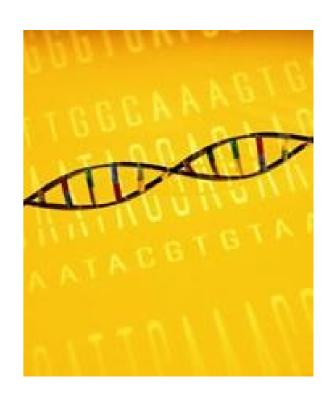


https://en.wikipedia.org/wiki/Silent\_mutation



#### Silent Mutations

- Are these mutations really so subtle?
- Are there dangers involved?
  - While the protein may be fine, the RNA has still has dangerous folding issues
- Nature: Silent Mutations Speak Up: Overlooked genetic changes could impact on disease
  - http://www.nature.com/news/2006 /061221/full/news061218-12.html





#### Second letter

			Michael Walter Control of Control		
	U	С	Α	G	
U	UUU } Phe UUA } Leu UUG }	UCU UCC UCA UCG	UAU Tyr UAC Stop UAG Stop	UGU Cys UGC Stop UGG Trp	U C A G
С	CUU CUC CUA CUG	CCU CCC CCA CCG	CAU His CAC Gln CAG	CGU CGC CGA CGG	U C A G
Α	AUU AUC AUA IIIe AUA Met	ACU ACC ACA ACG	AAU Asn AAA AAG Lys	AGU Ser AGA AGA AGG	U C A G
G	GUU GUC GUA GUG	GCU GCC GCA GCG	GAU Asp GAC Asp GAA GAA GAG	GGU GGC GGA GGG	U C A G

First letter

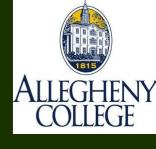
# Third letter



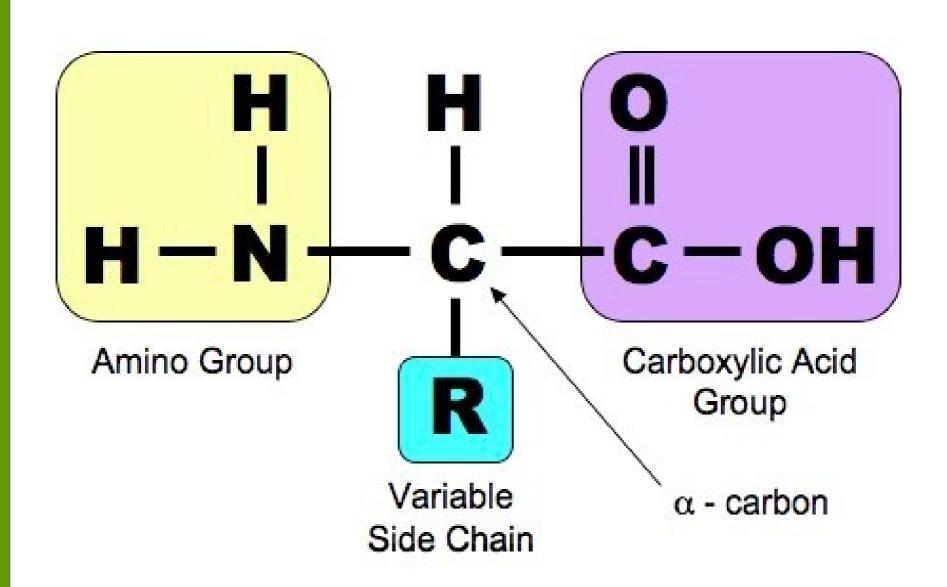
#### Alphabetical Interests

- With a larger protein "alphabet" (20 amino acids), it is much less likely to get matches by chance.
- Matches are likely to be statistically significance
- Amino acid changes are not equally harmful to protein structure
  - Chemical complexes being replaced by similar chemical complex.
  - Ex: Arginine (Arg) and Lysine (Lys)

$$H_2N$$
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 $H_2N$ 
 $H_2N$ 
 $H_2N$ 
 $H_2N$ 
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 $H_2N$ 
 $H_3$ 
 $H_3$ 



#### **Amino Acid Substitutions**







$\begin{array}{c c} H & O \\ & \parallel \\ H_2N-C-C-OH \\ & \parallel \\ H \\ \text{Glycine (Gly)} \end{array}$	H O        H <sub>2</sub> N-C-C-OH   CH <sub>3</sub> Alanine (Ala)			
O    N <sup>+</sup> C O H   H Proline (Pro)		$H$ O $\parallel$	H O      H2N—C—C—OH   CH2   OH   Serine (Ser)	H O      H <sub>2</sub> N-C-C-OH   HO-CH   CH <sub>3</sub>   Threonine (Thr)
OH Aspartic acid (Asp)	$\begin{array}{c c} H & O \\ & \parallel & \parallel \\ H_2N-C-C-OH \\ & CH_2 \\ & CH_2 \\ & C=O \\ & OH \\ & Glutamic acid (Glu) \end{array}$	H O      H <sub>2</sub> N-C-C-OH   CH <sub>2</sub>   C=O   NH <sub>2</sub> Asparagine (Asn)	$\begin{array}{c c} H & O \\ & \parallel & \parallel \\ H_2N-C-C-C-OH \\ & CH_2 \\ & CH_2 \\ & C=O \\ & NH_2 \\ & Glutamine (Glu) \end{array}$	$\begin{array}{c} H  O \\ \mid  \parallel \\ H_2N-C-C-OH \\ \mid  \\ CH_2 \\ \mid  \\ CH_2 \\ \mid  \\ CH_2 \\ \mid  \\ CH_2 \\ \mid  \\ \downarrow  \\ CH_2 \\ \mid  \\ Lysine (Lys) \end{array}$
H O             H <sub>2</sub> N-C-C-OH   CH <sub>2</sub>   CH <sub>2</sub>   CH <sub>2</sub>   NH   C=NH   NH <sub>2</sub>   Arginine (Arg)	$H_2N$ — $C$ — $C$ — $C$ — $OH$ $CH_2$ $N$ $H$ $H$ Histidine (His)	H O	H O           H <sub>2</sub> N-C-C-OH   CH <sub>2</sub>   OH   Tyrosine (Tyr)	$H$ O $H_2N-C-C-OH$ $CH_2$ $N$ $H$ Tryptophan (Trp)

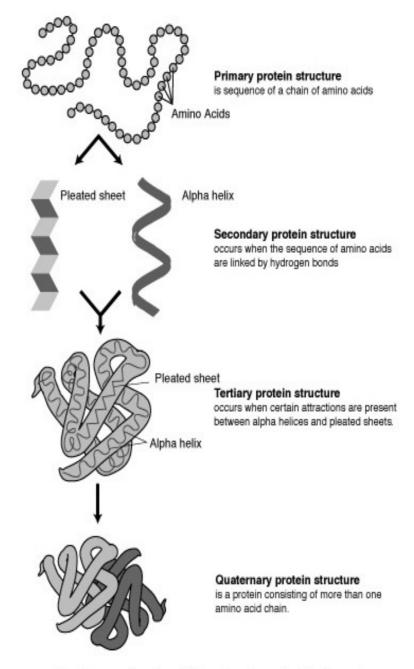


#### **Amino Acid Components**

- Similarity of amino acids means
  - Similar physicochemical properties (Physics + chemistry)
    - Polar vs nonpolar
    - Hydrophobic vs hydrophilic
    - Positive electric charge vs negative electric charge
    - Basic vs Acidic
- Amino Acid Table: <a href="http://www.bio.davidson.edu/courses/genomics/jmol/aatable.html">http://www.bio.davidson.edu/courses/genomics/jmol/aatable.html</a>
- Roles in Protein Structures
- http://www.proteinstructures.com/Structure/Structure/amino-acids.html



# Amino Acids Determine Protein's Shape and Function



The hierarchy of protein structure. Public domain image from The National Genome Research Institute



#### Scoring Amino Acid Substitutions

 Could we quantify sequence by physicochemical properties? (yes!)

**Table 5.1** Hydrophobicity values for the 20 amino acids. A more positive value represents a more hydrophobic amino acid.

Amino Acid	Hydrophobicity	Amino Acid	Hydrophobicity	Amino Acid	Hydrophobicity
D	-3.5	Υ	-1.3	1	4.5
K	-3.9	N	-3.5	С	2.5
Н	-3.2	L	3.8	А	1.8
Т	-0.7	E	-3.5	S	-0.8
V	4.2	R	-4.5	G	-0.4
F	2.8	W	-0.9	Р	-1.6
М	1.9	Q	-3.5		



#### Scoring Amino Acid Substitutions

Better to study evolution of real proteins from <u>closely</u> <u>related</u> organisms

Minimizes likelihood that an observed difference represents a series of more than one individual mutations

Species A – Ala

Species B – Ile

No intermediate mutations?

Ala --> Ile : 1 mutation

Ala --> Pro --> Ser --> Ile : 3 mutations

A few intermediate mutations?



## A Model of Evolutionary Change in Proteins, Dayhoff et al., 1978

Global Pairwise Alignment

Observed frequency of each possible amino acid substitution:

$$10 \log_{10} (M_{ij}/f_j)$$

- M<sub>ij</sub> the probability of a mutation replacing amino i with j
- f<sub>j</sub> the frequency of amino acid j in a large set of sequences



## A Model of Evolutionary Change in Proteins, Dayhoff et al., 1978

Global Pairwise Alignment

Observed frequency of each possible amino acid substitution:

$$10 \log_{10} (M_{ij}/f_i)$$

#### **Odds** ratio

- = 1 substitution of *j* for *i* is no more likely than the chance of finding *j* randomly
- > 1 substitution is evolutionarily conserved
- < 1 substitution is selected against



## A Model of Evolutionary Change in Proteins, Dayhoff et al., 1978

Global Pairwise Alignment

Observed frequency of each possible amino acid substitution:

 $10 \log_{10} \left( M_{ij} / f_j \right)$ 

log-odds ratio – easier for scoring

Greater positive for likely (conservative) substitutions

Greater negative for unlikely (non-conservative) substitutions

Multiplied by 10 and rounded to nearest integer

## The PAM Matrix

		Α	R	N	D	С	Q	Е	G	Н	1	L	K	М	F	Р	S	Т	W	Υ	٧
Ala	Α	2																			
Arg	R	-1	5																		
Asn	N	0	0	3																	
Asp	D	0	-1	2	5																
Cys	С	-1	-1	-1	-3	11															
Gln	Q	-1	2	0	1	-3	5														
Glu	Е	-1	0	1	4	-4	2	5													
Gly	G	1	0	0	1	-1	-1	0	5												
His	Н	-2	2	1	0	0	2	0	-2	6											
lle	1	0	-3	-2	-3	-2	-3	-3	-3	-3	4										
Leu	L	-1	-3	-3	-4	-3	-2	-4	-4	-2	2	5			(						
Lys	K	-1	4	1	0	-3	2	1	-1	1	-3	-3	5								
Met	М	-1	-2	-2	-3	-2	-2	3	3	-2	3	3	-2	6							
Phe	F	-3	-4	-3	<b>-</b> 5	0	-4	<b>-</b> 5	-5	0	0	2	-5	0	8						
Pro	Р	1	-1	-1	-2	-2	0	-2	-1	0	-2	0	-2	-2	-3	6					
Ser	S	1	-1	1	0	1	-1	-1	1	-1	-1	-2	-1	-1	-2	1	2				
Thr	Т	2	-1	1	-1	-1	-1	-1	-1	-1	1	-1	-1	0	-2	1	1	2			
Trp	W	-4	0	-5	-5	1	-3	<b>-</b> 5	-2	-3	-4	-2	-3	-3	-1	-4	-3	-4	15		
Tyr	Υ	-3	-2	-1	-2	2	-2	-4	-4	4	-2	-1	-3	-2	5	-3	-1	-3	0	9	
Val	٧	1	-3	-2	-2	-2	-3	-2	-2	-3	4	2	-3	2	0	-1	-1	0	-3	-3	4



#### PAM matrices

- Point Accepted Mutation
- Family of matrices PAM 1, PAM 80, PAM 120, PAM 250
- The number in the name of a PAM matrix (i.e., the 'n' in PAM n) represents the evolutionary distance between the sequences on which the matrix is based

BLOSUM 80

PAM 1

PAM 120

BLOSUM 45

PAM 250

Less divergent

More divergent



#### PAM vs BLOSUM

- General Use
  - PAM 120
  - BLOSUM 62\*
- Closely Related Species
  - PAM 60
  - BLOSUM 80
- Distantly Related Species
  - PAM 250
  - BLOSUM 45

PAM	BLOSUM
PAM100	BLOSUM90
PAM120	BLOSUM80
PAM160	BLOSUM60
PAM200	BLOSUM52
PAM250	BLOSUM45

\*BLOSUM 62 – used by BLAST – computed by choosing blocks of local alignments more than 62% identical



#### **Blast Subst Matrices**

- Scoring for possible residue pair alignment
- Different substitution matrices are for detecting similarities according to degrees of divergence.
- BLOSUM-62 matrix good for detecting most weak protein similarities
- Provisional table of recommended substitution matrices and gap costs for various query lengths is

Query Length	Substitution Matrix	Gap Costs
<35	PAM-30	(9,1)
35-50	PAM-70	(10,1)
50-85	BLOSUM-80	(10,1)
85	BLOSUM-62	(10,1)



#### BLOSUM matrix Heinkoff and Heinkoff, 1992

BLOcks SUbstition Matrix - Blocks of local alignments

$$S_{ij} = \left(\frac{1}{\lambda}\right) \log \left(\frac{p_{ij}}{q_i * q_j}\right)$$

- p<sub>ii</sub> probability j replacing i
- $q_i$  and  $q_j$  probabilities of finding the amino acids i and j in any protein sequence
- λ scaling factor, set such that the matrix contains easily computable integer values.
- BLOSUM # # = minimum % similarity of sequences compared





- Create N x M matrix
- Place each sequence along one axis
- Place score 0 at the up-left corner
- Fill in 1<sup>st</sup> row & column with gap penalty multiples
- Fill in the matrix with max value of 3 possible moves:
  - Vertical move: Score + gap penalty
  - Horizontal move: Score + gap penalty
  - Diagonal move: Score + match/mismatch score
- The optimal alignment score is in the lower-right corner
- To reconstruct the optimal alignment, trace back where the max at each step came from, stop when hit the origin.

#### Needleman-Wunsch Algorithm: Protein Alignment – Chap 5



- Create N x M matrix
- Place each sequence along one axis
- Place score 0 at the up-left corner
- Fill in 1<sup>st</sup> row & column with gap penalty multiples
- Fill in the matrix with max value of 3 possible moves:
  - Vertical move: Score + gap penalty
  - Horizontal move: Score + gap penalty
  - Diagonal move: Score + match/mismatch score from sub. matrix
- The optimal alignment score is in the lower-right corner
- To reconstruct the optimal alignment, trace back where the max at each step came from, stop when hit the origin.

#### Blast-Off!!



- Let's blast some protein sequences
- https://blast.ncbi.nlm.nih.gov/Blast.cgi#dtr\_Quer y\_98931



