Proteins: Structure & Function

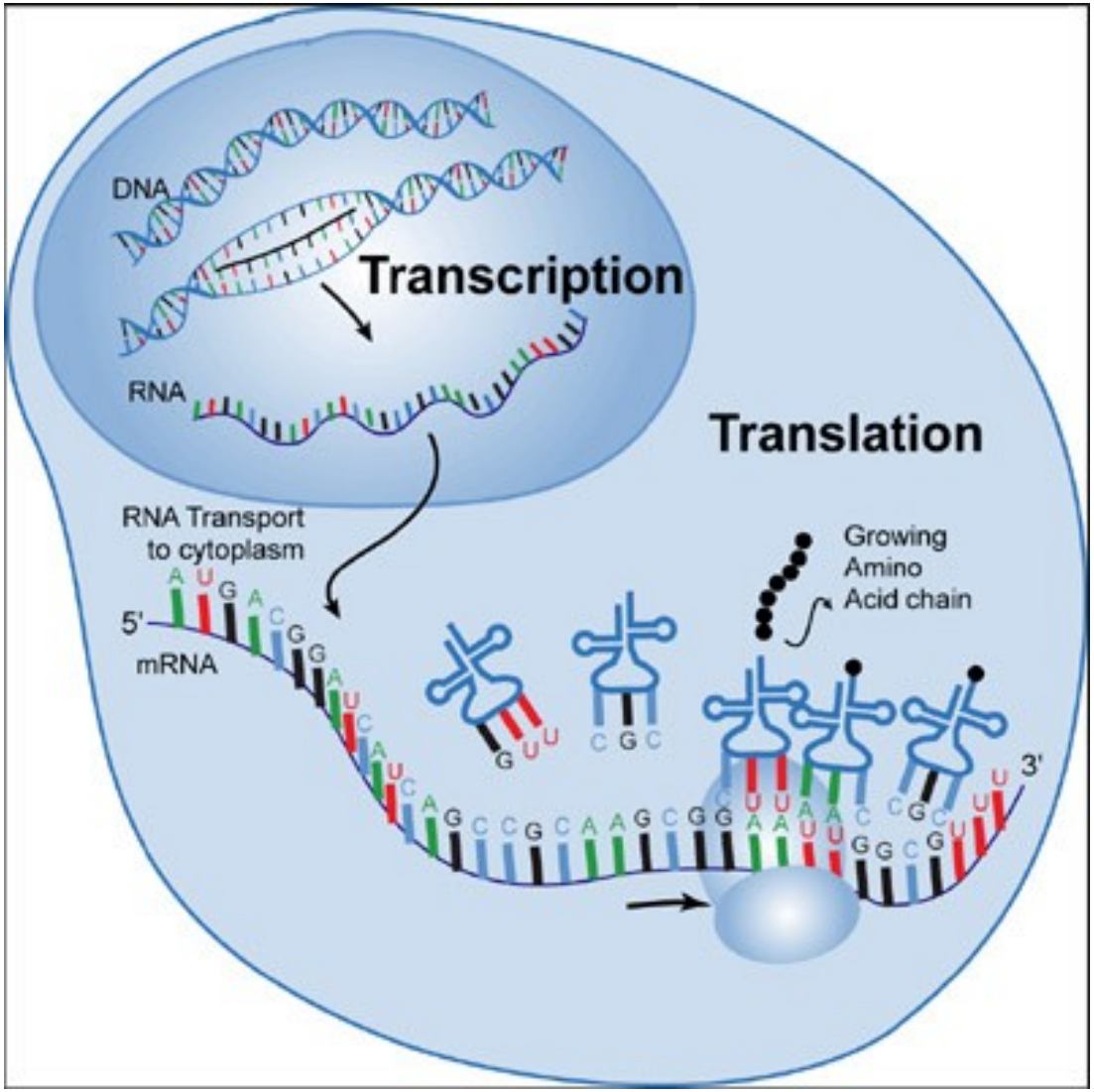
Johannes Starlinger

This Lecture

- Proteins
  - Structure
  - Function
  - Databases
- Predicting Protein Secondary Structure

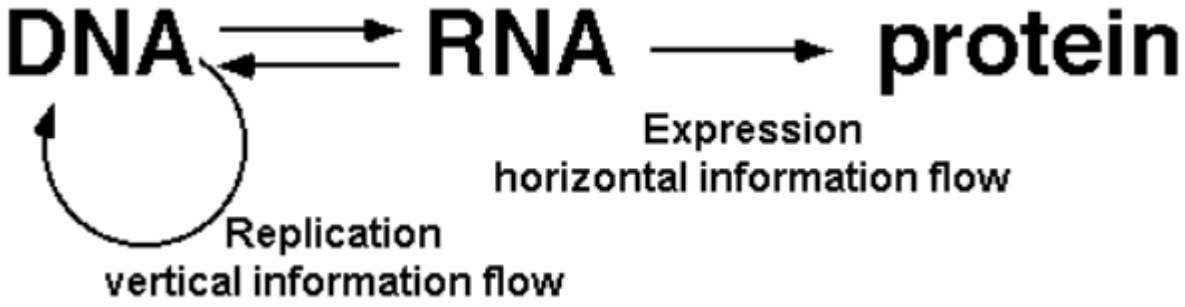
- Many figures from Zvelebil, M. and Baum, J. O. (2008). "Understanding Bioinformatics", Garland Science, Taylor & Francis Group.
- Examples often from O. Kohlbacher, Vorlesung Strukturvorhersage, WS 2004/2005, Universität Tübingen

DNA→**Transcription**→RNA→**Translation**→Protein



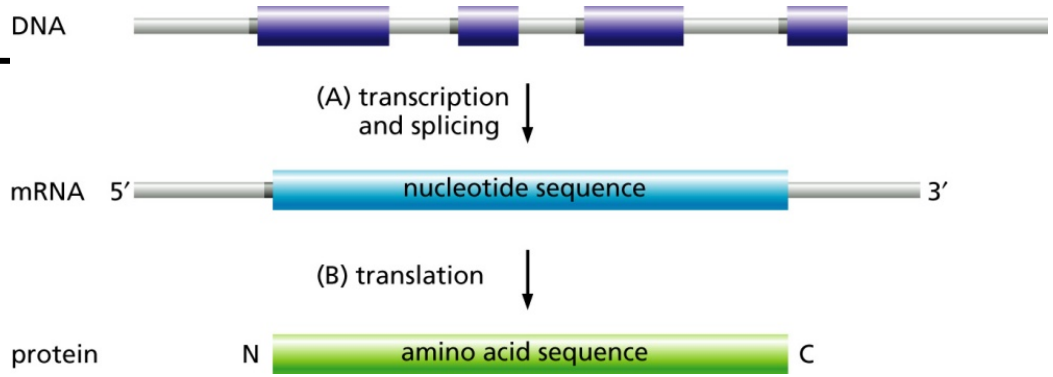
From:
http://www.tokresource.org

Central Dogma of Molecular Biology



	U		C		A		G	
U	UUU UUC	Phenyl-alanine	UCU UCC UCA UCG	Serine	UAU UAC	Tyrosine	UGU UGC	Cysteine	U C A G
	UUA UUG	Leucine			UAA UAG	Stop codon Stop codon	UGA	Stop codon
							UGG	Tryptophan
C	CUU CUC CUA CUG	Leucine	CCU CCC CCA CCG	Proline	CAU CAC	Histidine	CGU CGC CGA CGG	Arginine	U C A G
					CAA CAG	Glutamine		
A	AUU AUC AUA	Isoleucine	ACU ACC ACA ACG	Threonine	AAU AAC	Asparagine	AGU AGC	Serine	U C A G
	AUG	Methionine; initiation codon			AAA AAG	Lysine	AGA AGG	Arginine
G	GUU GUC GUA GUG	Valine	GCU GCC GCA GCG	Alanine	GAU GAC	Aspartic acid	GGU GGC GGA GGG	Glycine	U C A G
					GAA GAG	Glutamic acid		

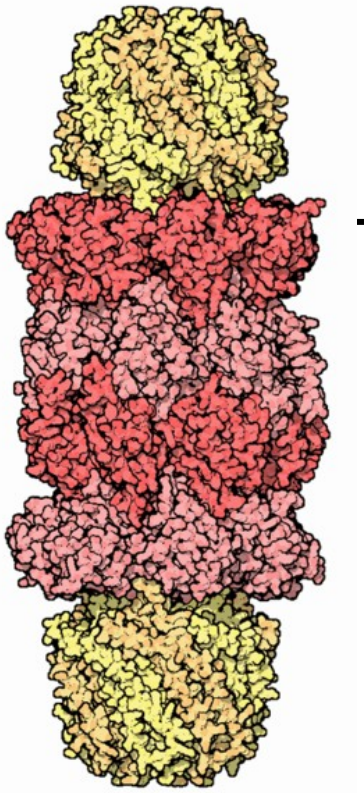
Details

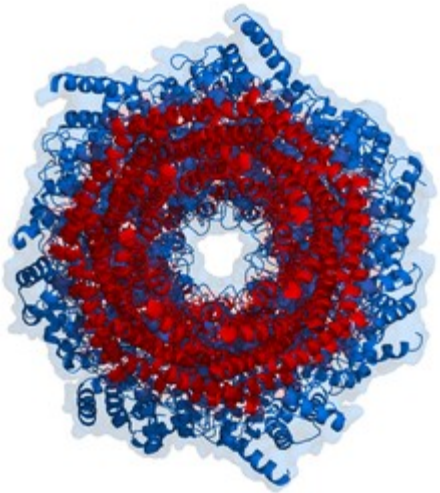


- Alternative Splicing
  - "One gene – one protein" is wrong
  - Exons may be spliced out from the mRNA
  - Human: ~6 times more different proteins than genes
- Post-translational modifications
  - (De-)Phosporylation, glycolysation, cleavage of signals, …
  - Rough estimates: 100K proteins, 500K protein forms
- Complexes: Proteins physically group together to perform specific function

Example: Proteasome

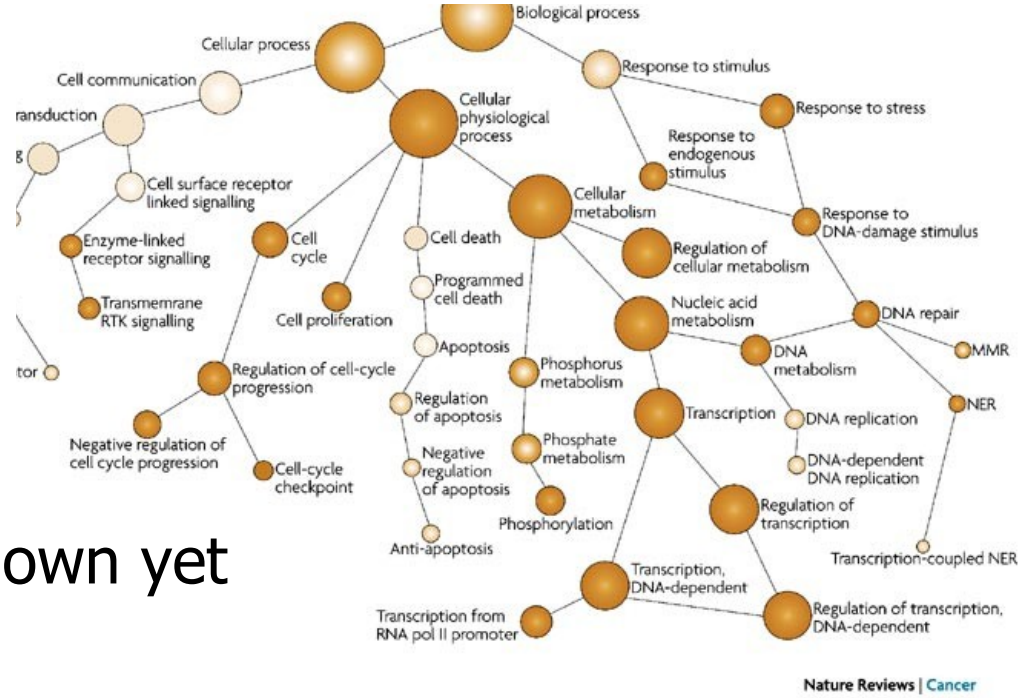
- Very large complexes present in all eukaryotes (and more species)
  - >2000 kDa, made of dozens of single protein chains
  - Formation of the complex is a complex process only partly understood yet
- Breaks (mis-folded, broken, superfluous, …) proteins into small peptides for reuse



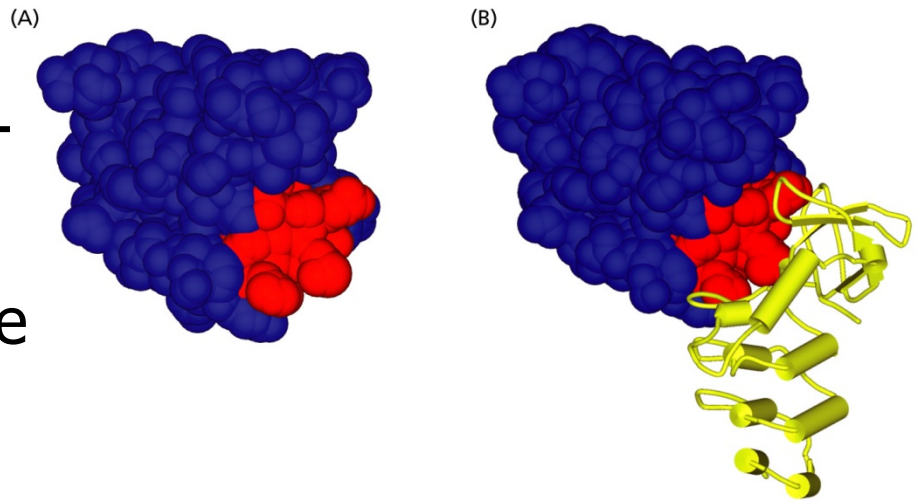


Protein Function

- Proteins perform essentially everything that makes an organism alive
  - Metabolism
  - Signal processing
  - Gene regulation
  - Cell cycle
  - …
- For ~1/3 of all human genes, no function is known yet
- Describing function
  - Gene Ontology: 3 branches, >30.000 concepts
    - Cellular component, biological process, molecular function
  - Used world-wide to describe gene/protein function



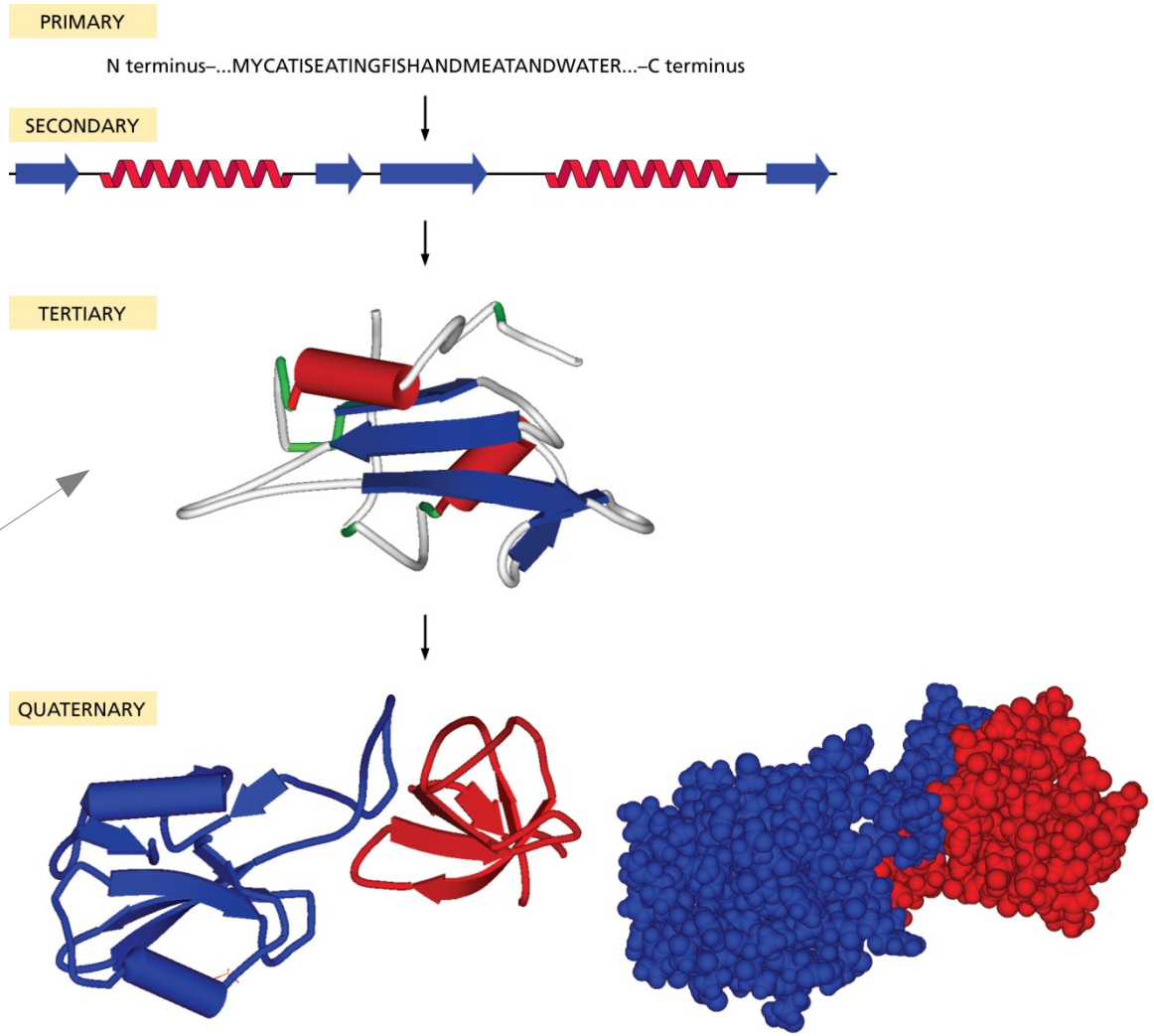
Function and Motifs



- Proteins often have multiple functions
  - Avg. no. of GO terms assigned to a human protein: ~6
- Functions are associated to motifs or domains
- There probably exist only 4000-5000 motifs
  - Proteins as assemblies of functional motifs
- Performing a function often requires binding to another protein or molecule
  - The binding requires a certain constellation of the protein structure
  - Major target of pharmacological research

Protein Structure

- Primary
  - 1D-Seq. of AA
- Secondary
  - 1D-Seq. of "subfolds"
- Tertiary
  - 3D-Structure
- Quaternary
  - Assembled complexes



Protein Structure

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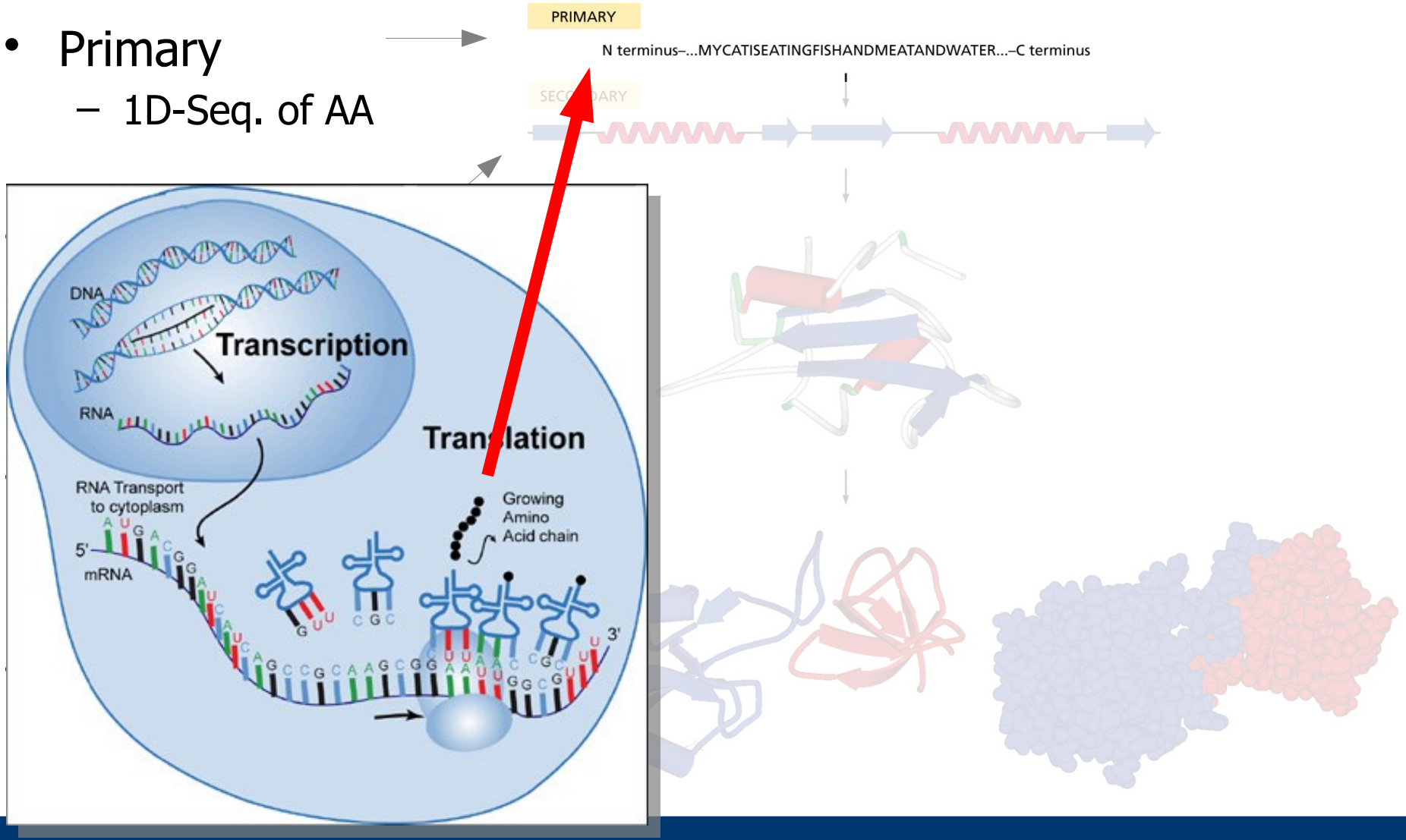
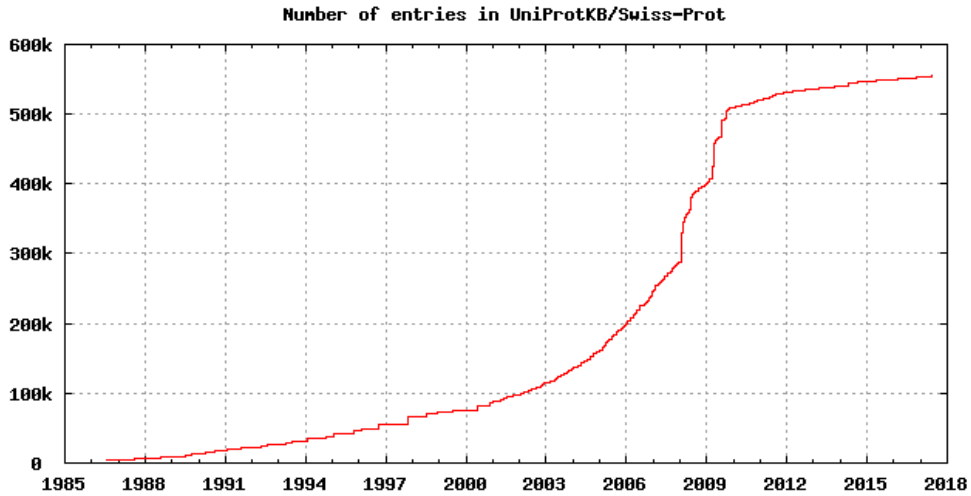
  - 1D-Seq. of AA



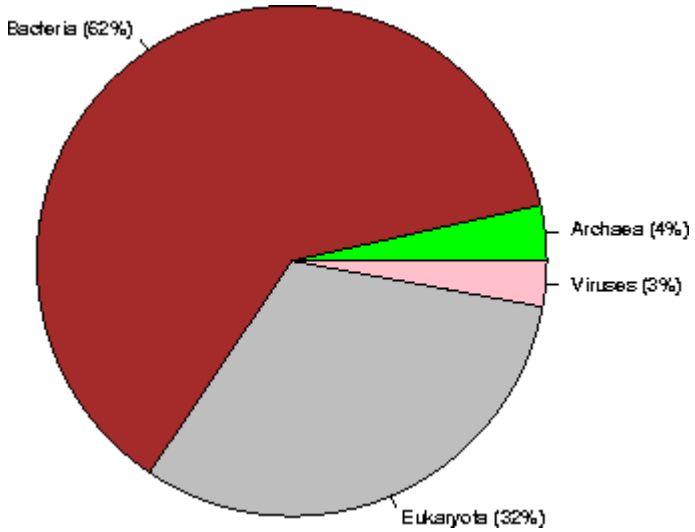
UniProt

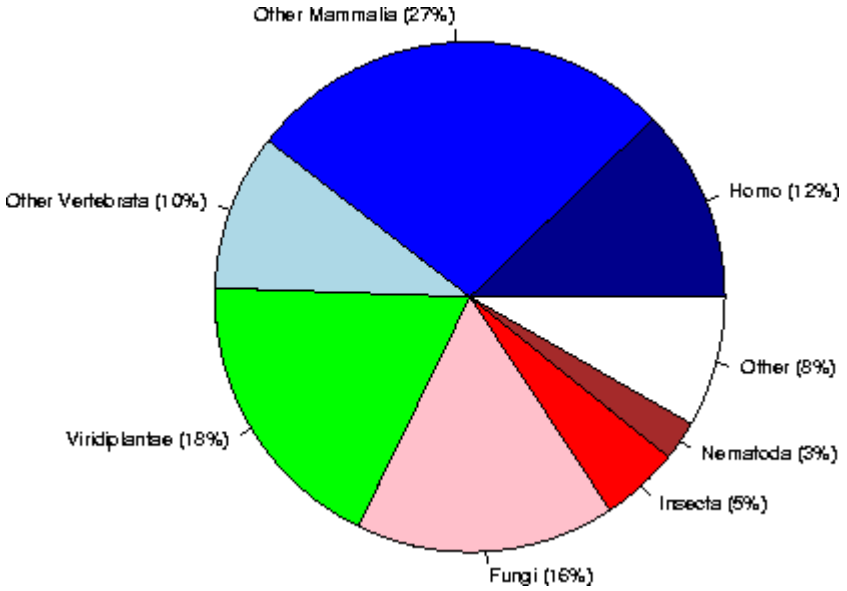
- "Standard" database for protein sequences and annotation
  - Original name: SwissProt
  - Started at the Swiss Institute of Bioinformatics, now mostly EBI
  - Other: PIR, HPRD
- Continuous growth and curation
  - >30 „Scientific Database Curators"
  - Quarterly releases
  - Very rich set of annotations



- Actually two databases
  - SwissProt: Curated, high quality, versioned
  - TrEMBL: Automatic generation from (putative) coding genomic sequences, low quality, redundant, much larger

UniProt: Species [http://www.expasy.org/sprot/relnotes/relstat.html]

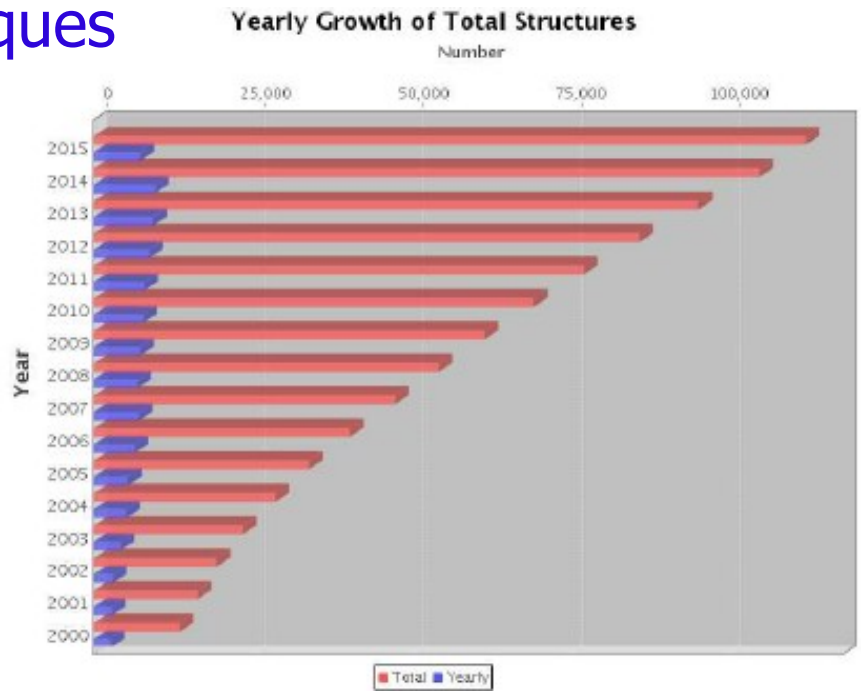




Number	Frequency	Species
1	20205	Homo sapiens (Human)
2	16894	Mus musculus (Mouse)
3	15383	Arabidopsis thaliana (Mouse-ear cress)
4	7991	Rattus norvegicus (Rat)
5	6721	Saccharomyces cerevisiae (Baker's yeast)
6	5999	Bos taurus (Bovine)
		...

PDB – Protein Structure Database

- Oldest protein database, evolved from a book
- Contains experimentally obtained protein 3D-structures
  - Plus some DNA, protein-ligand, complexes, …
  - X-Ray (~75%), NMR (nuclear magnetic resonance, ~23%)
- Costly and rather slow techniques
  - Growth much smaller than that of sequence-related DBs
- Many problems with legacy data and data formats

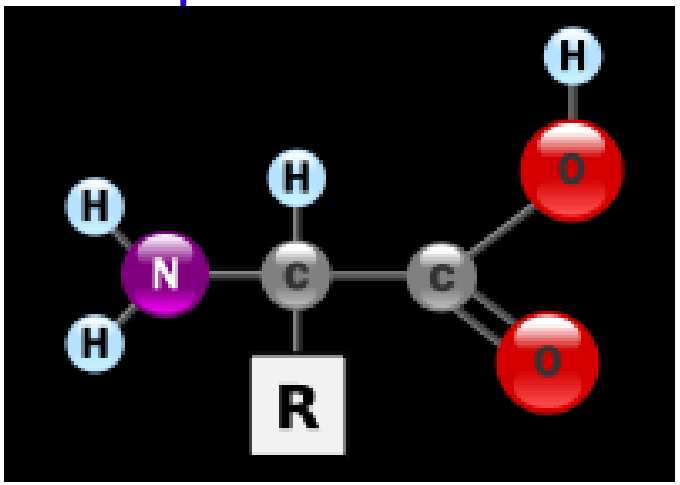


This Lecture

- Introduction
- Predicting Protein Secondary Structure
  - Secondary structure elements
  - Chou-Fasman
  - Other methods

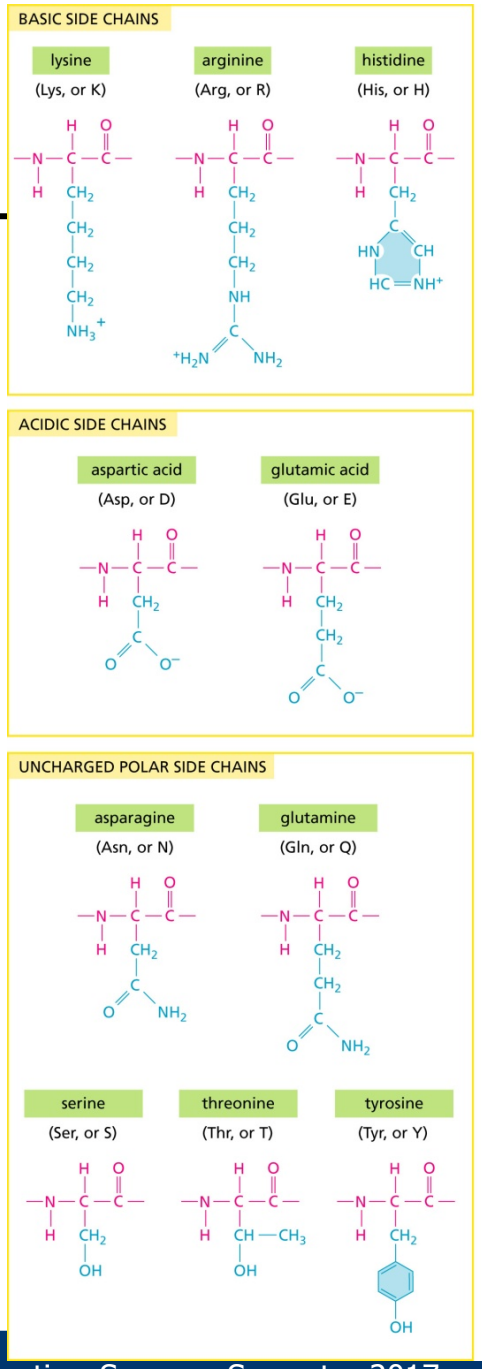
Amino Acids

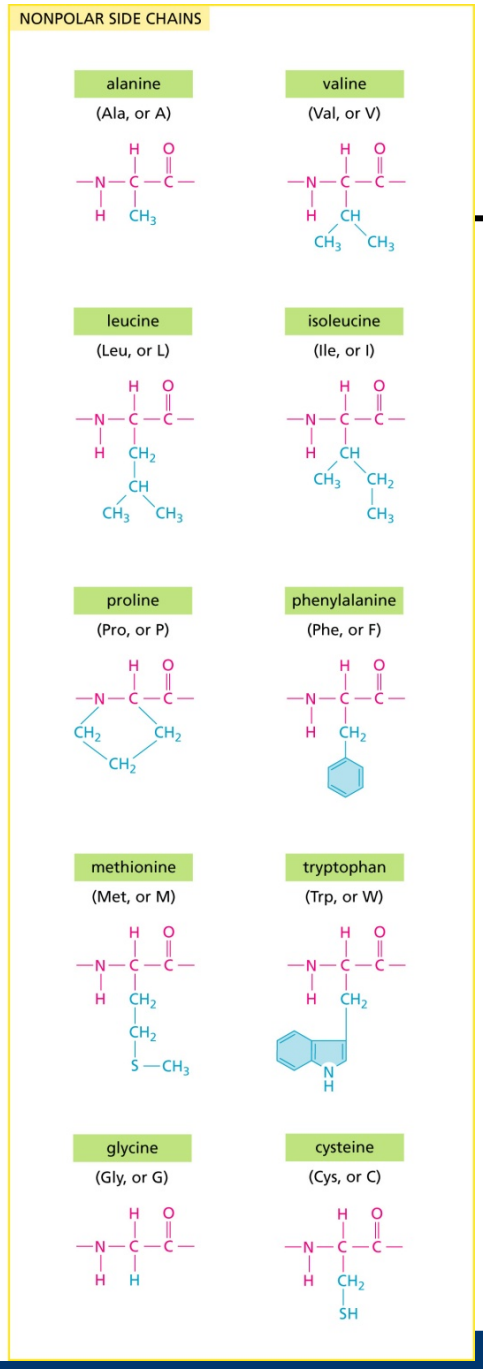
- AA consist of a common core and a specific residue
  - Amino group – NH_2
  - Central C_α - Carbon – CH
  - Carboxyl group – COOH



- Residues (side chains) vary greatly between AA
- Residues determine the specific properties of a AA

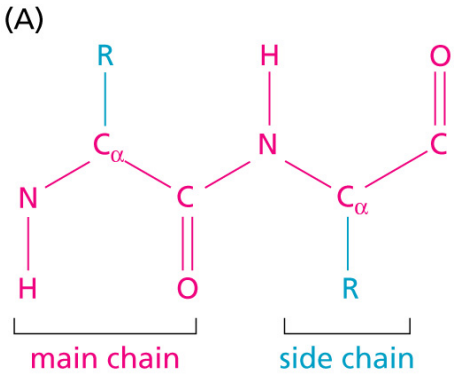
Side Chains

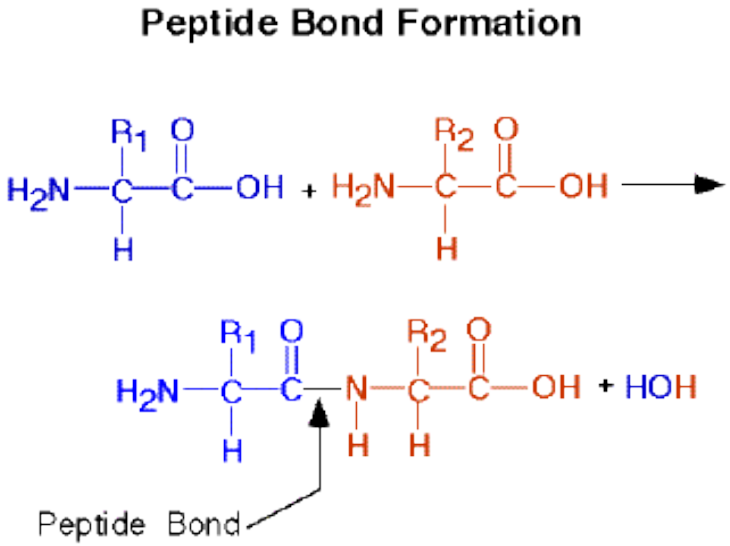

BASIC SIDE CHAINS
lysine
(Lys, or K)
arginine
(Arg, or R)
histidine
(His, or H)
ACIDIC SIDE CHAINS
aspartic acid
(Asp, or D)
glutamic acid
(Glu, or E)
UNCHARGED POLAR SIDE CHAINS
asparagine
(Asn, or N)
glutamine
(Gln, or Q)
serine
(Ser, or S)
threonine
(Thr, or T)
tyrosine
(Tyr, or Y)


NONPOLAR SIDE CHAINS
alanine
(Ala, or A)
valine
(Val, or V)
leucine
(Leu, or L)
isoleucine
(Ile, or I)
proline
(Pro, or P)
phenylalanine
(Phe, or F)
methionine
(Met, or M)
tryptophan
(Trp, or W)
glycine
(Gly, or G)
cysteine
(Cys, or C)

Structure of a Protein

- Concatenation of cores: Backbone of AA chain (a protein)
- Covalent peptide bonds between carboxyl and amino group (with loss of H_2O)





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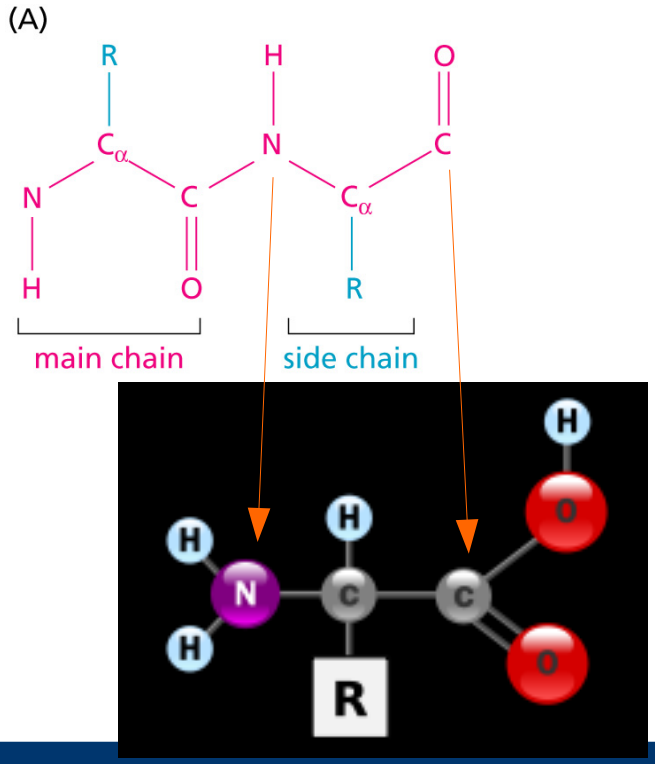

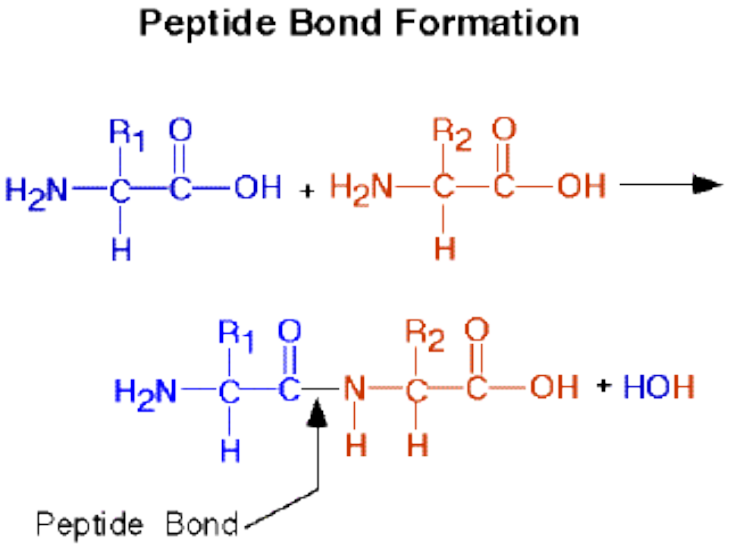

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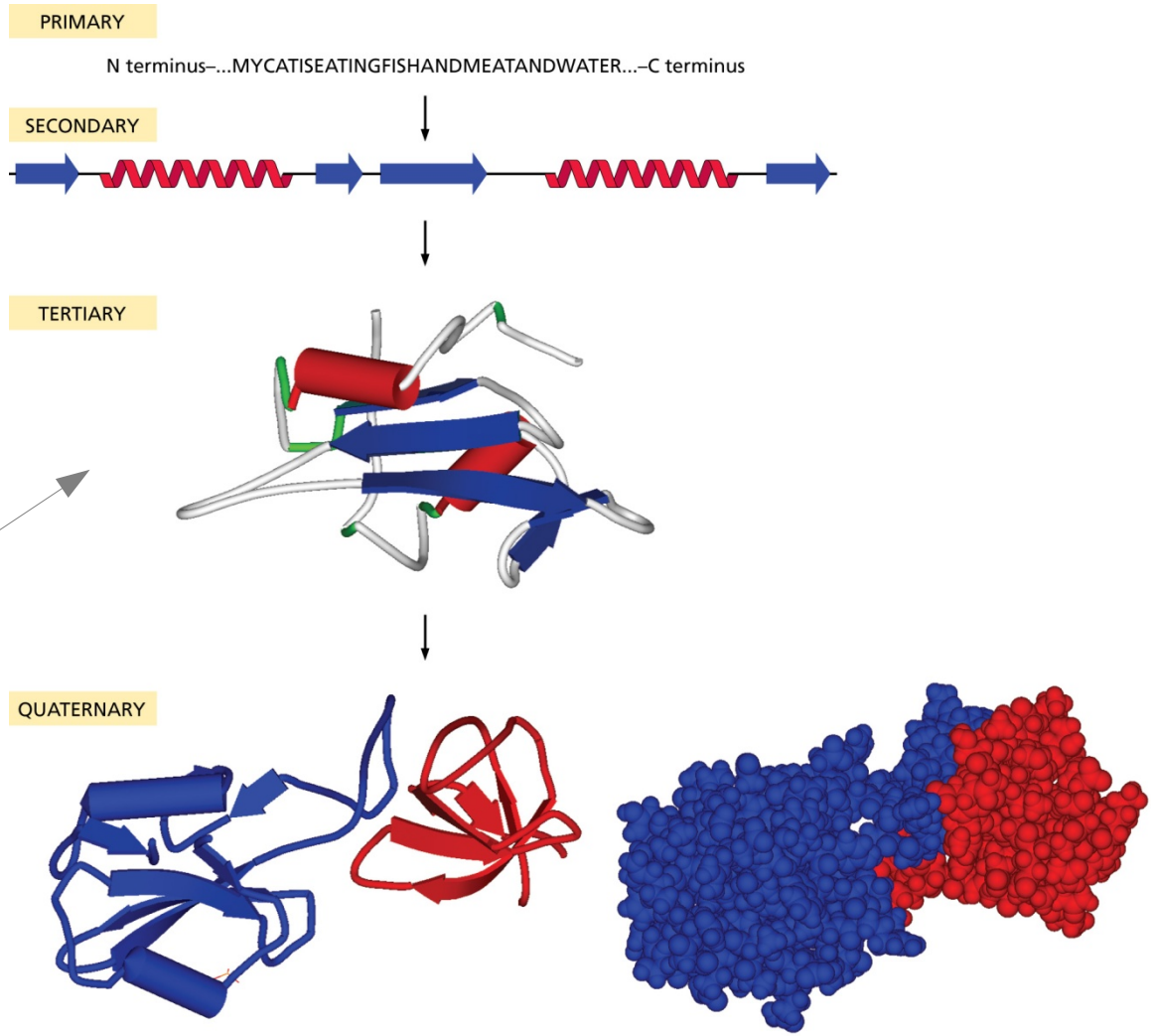
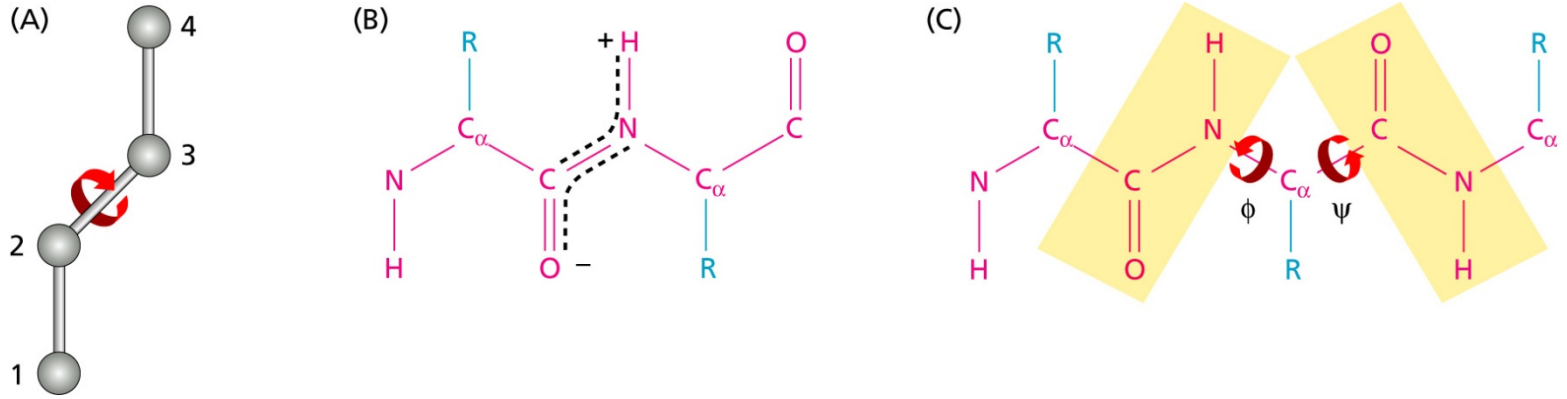


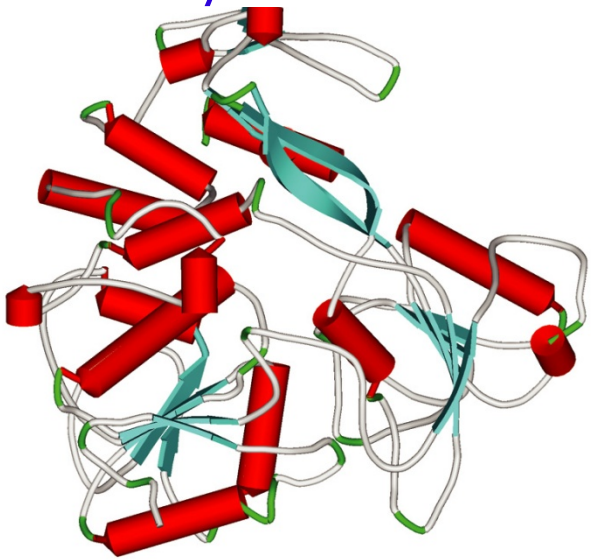
Flexibility

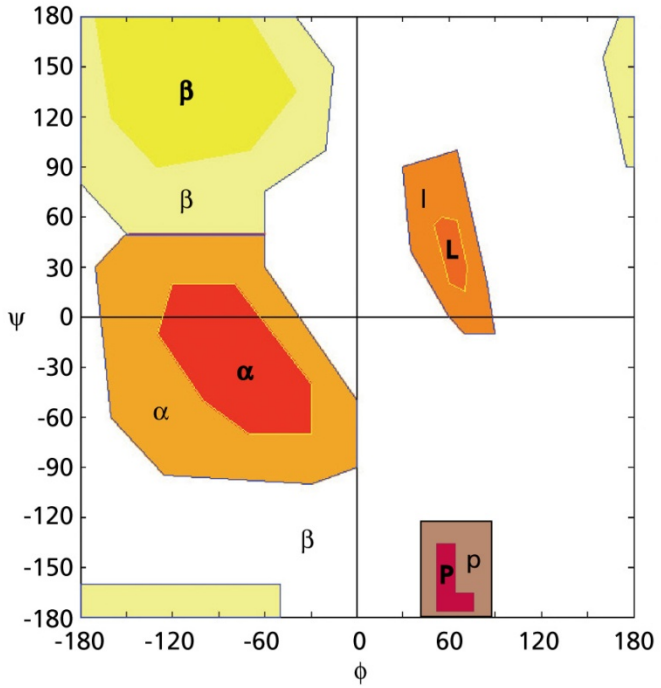
- In principle, every chemical bond can rotate freely
- Would allow arbitrary backbone structures
- In proteins things are more restricted
  - Peptide bound is "flat" – almost no torsion possible
  - Flexibility only in the C_α-flanking bonds ϕ and ψ



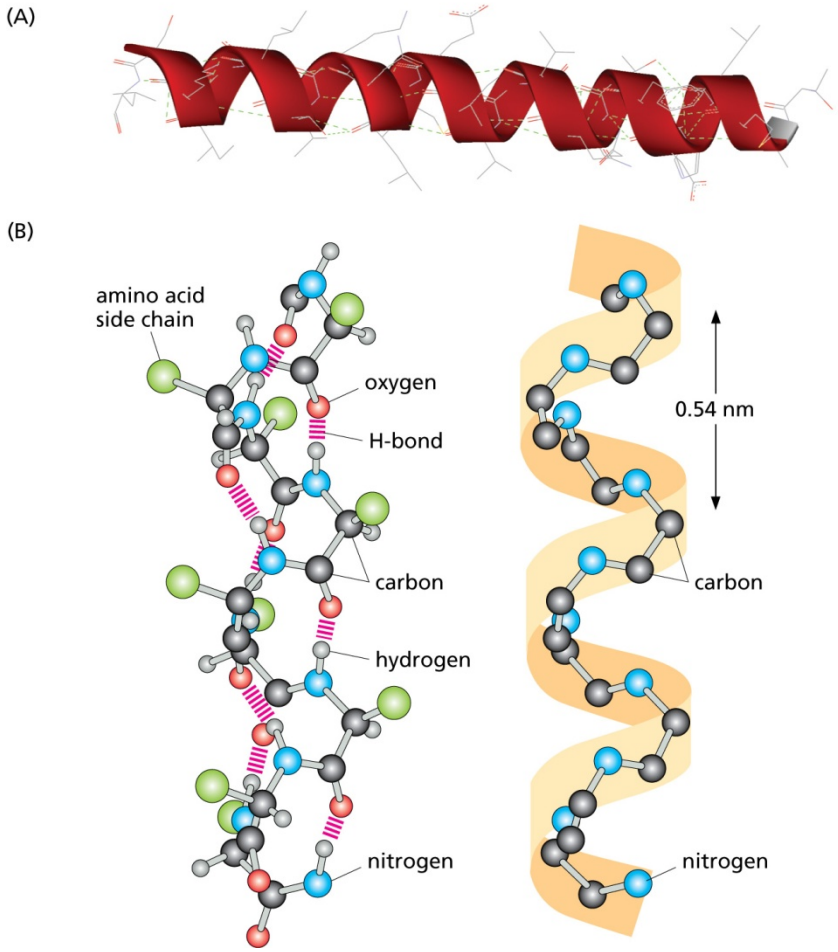
Ramachandran Plots

- Combinations of ϕ and ψ are highly constrained
  - Due to chemical properties of the backbone / side chains
- Two combinations are favored:
  α-helixes and β-sheets
  - More detailed classifications exist
  - Angles lead to specific structures
  - Secondary structure



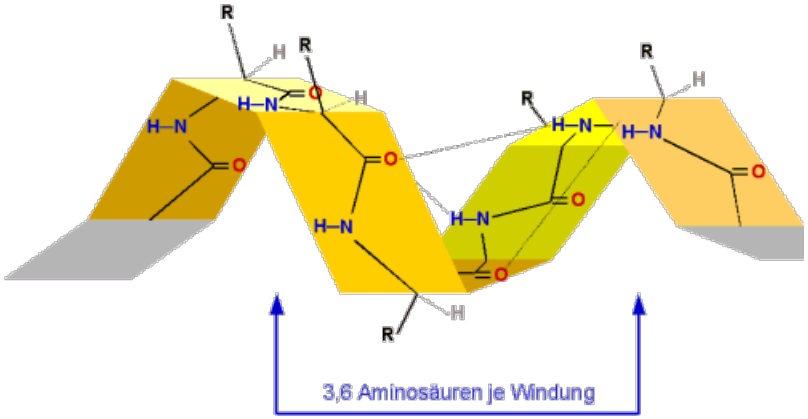


α-Helix



- Sequence of angles forming a regularly structured helix
- Additional bonds between amino and carboxyl groups
  - Very stable structure
- May have two orientations
  - Most are right-handed
- 3.4 AA per twist
- Often short, sometimes very long

α-Helix



From:
www2.chemie.uni-erlangen.de

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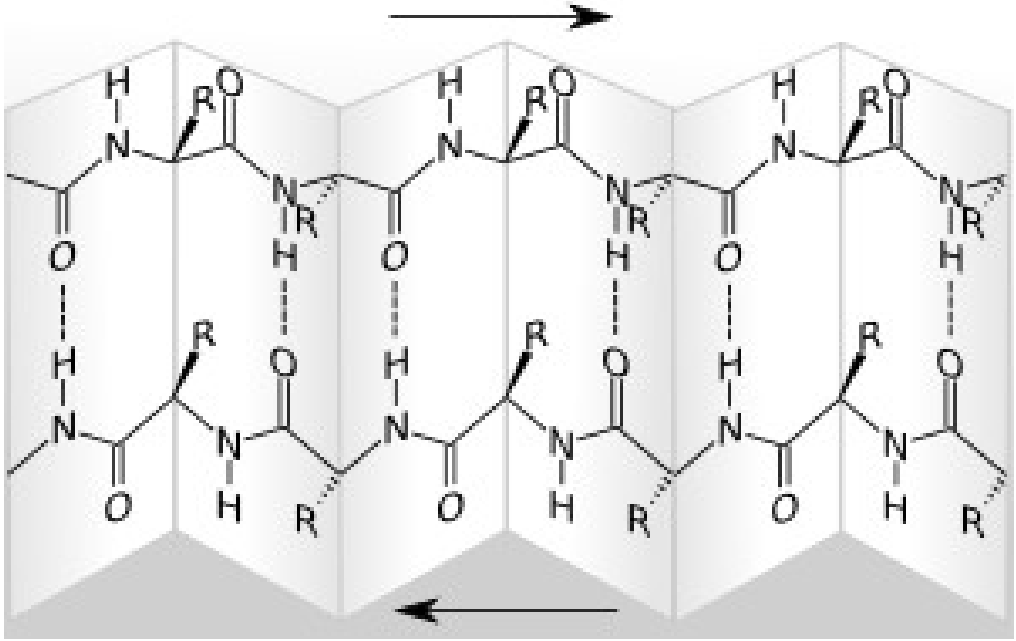
β-Sheet

- Two linear and parallel stretches (β-strands)
- Strands are bound together by hydrogen bounds
- Can be parallel or anti-parallel (wrt. N/C terminus)



Quelle: Wikipedia

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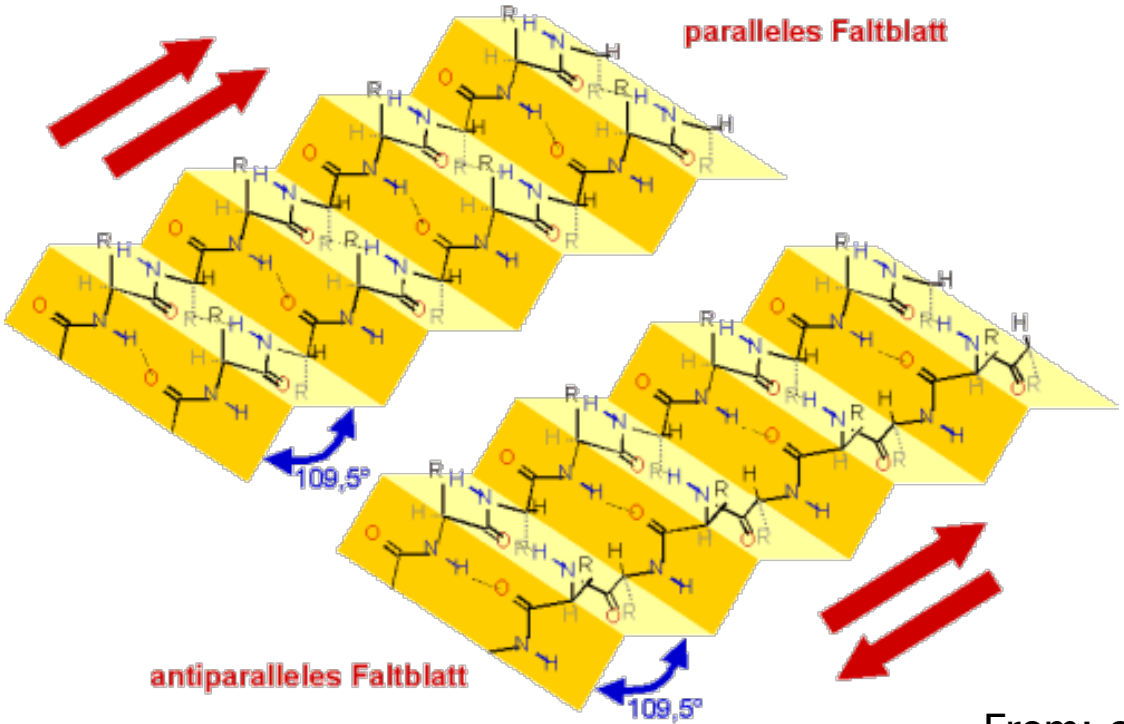
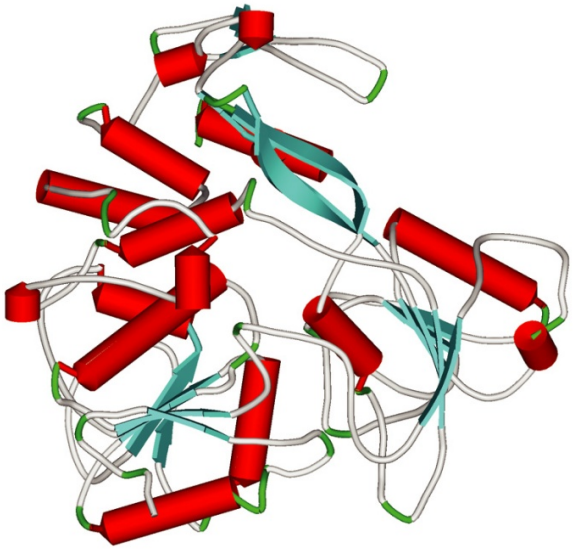


From: chemgapedia.de

Other Substructures

- α-helixes and β-sheets form around 50-80% of a protein
- Other parts are called loops or coils
  - Usually not very important for the structure of the protein
  - But very important for its function
  - Often exposed on the surface; important for binding to other molecules



Importance

- Secondary structure elements (SSE) are vital for the overall structure of a protein
- Often evolutionarily well conserved
- SSE can be used to classify proteins
  - Such classes are highly associated to function
- Knowing the SSE gives important clues to protein function
- Secondary structure prediction (SSP) is much simpler than 3D structure prediction
  - And 3D structure prediction can benefit a lot from a good SSP

Predicting Secondary Structure

- SSP: Given a protein sequence, assign each AA in the sequence to one of the three classes Helix (H), Strand (E), or Coil (-)

```
KVYGRCELAAAMKRLGLDNYRGYSLGNWVCAAKFESNFNTHATNRNTD
GSTDYGILQINSRWWCNDGRTPGSKNLCNIPCSALLSSDITASVNCAK
KIASGGNGMNAWVAWRNRCKGTDVHAWIRGCRL
```

```
KVYGRCELAAAMKRLGLDNYRGYSLGNWVCAAKFESNFNTHATNRNTD
-----HHHHHHHHH-------------EEEEE----------------
GSTDYGILQINSRWWCNDGRTPGSKNLCNIPCSALLSSDITASVNCAK
----EEEEEE-------------------------------HHHHHH
KIASGGNGMNAWVAWRNRCKGTDVHAWIRGCRL
HHH-------EEE--------------------
```

Classification

- Classification: Classify each AA into one of three classes
- Classification is a fundamental problem
  - Classify the readout of a microarray as diseased / healthy
  - Classify a subsequence of a genome as coding / non-coding
  - Classify an email as spam / no spam
- Many different techniques: Naïve Bayes, Regression, Decision Trees, SVMs, Neural Networks, …
  - Based on same principles can be exchanged easily
  - Classification function learned from properties of known objects
- The following is a rather unsystematic approach
  - But simple to explain and classical for this application

This Lecture

- Introduction
- Predicting Protein Secondary Structure
  - Secondary structure elements
  - Chou-Fasman
  - Other methods

Chou-Fasmann Algorithm

Chou & Fasman (1974). Prediction of protein conformation. Biochemistry 13

- Observation: Different AA favor different folds
  - Different AA are more or less often in H, E, C
  - Different AA are more or less often within, starting, or ending a stretch of H, E, C
- Chou-Fasman algorithm (rough idea)
  - Classifies each AA into E or H; unclassified AA are assigned C
  - Compute a score for the probability of any AA to be E (H)
  - Basis: Relative frequencies in a set of sequences with known SSE
  - In principle, assigns each AA its most frequent class
  - Add constraints about minimal length of E (H) stretches
  - Several further heuristics

Some Details [sketch, some heuristics omitted]

- Let $f_{j,k}$ be the relative frequency of observing AA j in class k
- Let f_k be the average over all 20 $f_{j,k}$ values
- Compute the propensity of AA j to be part of class k as

$$P_{j,k}=f_{j,k}/f_k$$

- Using $P_{j,k}$, classify each AA j for every class k into
  - Strong, normal, weak builder (H_α, h_α, I_α)
  - Strong, weak breaker (B_α, b_α)
  - Indifferent (i_α)

Concrete Values

- Originally computed on only 15 proteins (1974)

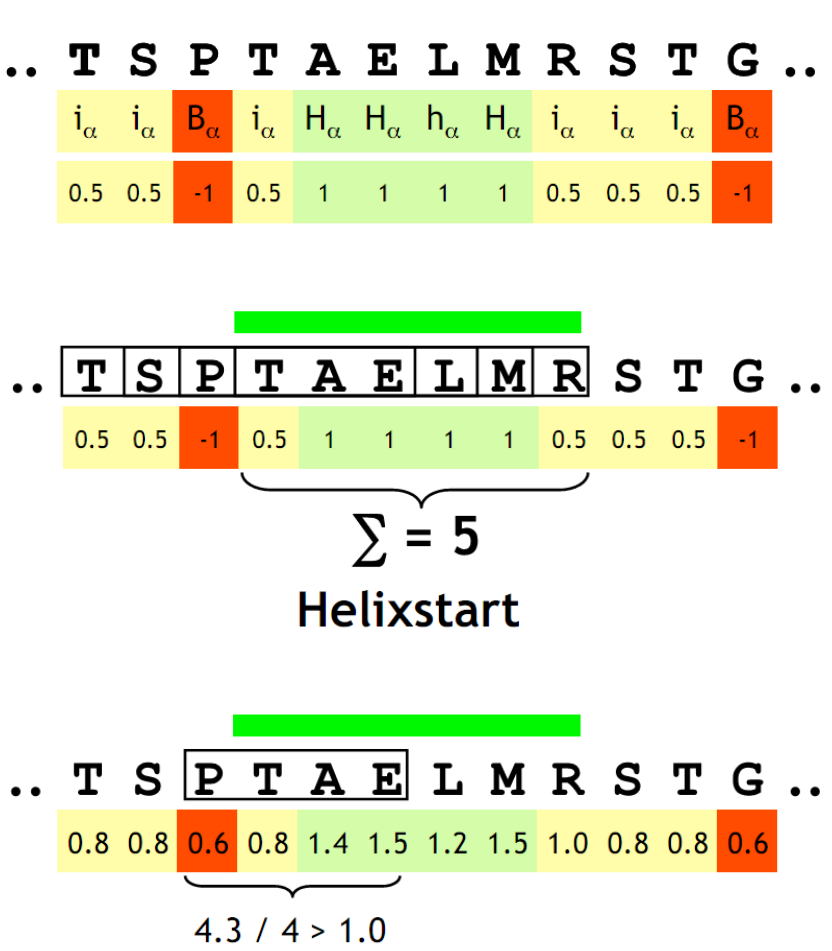
AS	P_α	Klasse	AS	P_β	Klasse
Glu	.53		Met	.67	
Ala	1.45	H_α	Val	1.65	H_β
Leu	1.34		Ile	1.60	
His	1.24		Cys	1.30	
Met	.20		Tyr	1.29	
Gln	1.17	h_α	Phe	1.28	h_β
Trp	1.14		Gln	1.23	
Val	1.14		Leu	1.22	
Phe	1.12		Thr	1.20	
Lys	1.07	I_α	Trp	1.19	

AS	P_α	Klasse	AS	P_β	Klasse
Ile	1.00	I_α	Ala	0.93	I_β
Asp	0.98		Arg	0.90	
Thr	0.82		Gly	0.81	i_β
Ser	0.79	i_α	Asp	0.80	
Arg	0.79		Lys	0.74	
Cys	0.77		Ser	0.72	
Asn	0.73	b_α	His	0.71	b_β
Tyr	0.61		Asn	0.65	
Pro	0.59	B_α	Pro	0.62	
Gly	0.53		Glu	.26	B_β

Algorithm for Helices

- Go through the protein sequence
- Score each AA with 1 (H_α, h_α), 0.5 (I_α, i_α), or -1 (B_α, b_α)
  - based on known values from propensity table
- Find helix cores: subsequences of length 6 with an aggregated AA score ≥ 4
- Starting from the middle of each core, shift a window of length 4 to the left (then to the right)
  - Compute aggregated score A using values $P_{j,k}$ inside the window
  - If A ≥ 4, continue; otherwise stop
- Similar method for strands
- Conflicts (regions assigned both H and E) are resolved based on aggregated scores

Example [Source: O. Kohlbacher, "Strukturvorhersage"]



Performance

- Accuracy app. 50-60%
  - Measured on per-AA correctness
- Prediction is more accurate in helices than in strands
  - Because helices build local bridges (hydrogen bounds between the turns; each AA binds to the +4 AA)
- General problem
  - Secondary structure is not only a local problem
  - Looking only at single AAs is not enough
    - Note: Scores are based on individual AA; aggregation by summation assumes statistical independence of pairs, triples … in a class
- One needs to include the context of an AA

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- Predicting Protein Secondary Structure
  - Secondary structure elements
  - Chou-Fasman
  - Other methods

Classes of Methods

- First generation: Properties of single AA only
  - Accuracy: 50-60%
  - E.g. Chou-Fasman (1974)
- Second generation: Include info. about neighborhood
  - Accuracy: ~65%
  - E.g. GOR (1974 – 1987)
- Third generation: Include info. from homologous seq's
  - Accuracy: ~70-75%
  - E.g. PHD (1994)
- Forth generation: Build ensembles of good methods
  - Accuracy: ~80%
  - E.g. Jpred (1998)

Further Reading

- Gerhard Steger (2003). “Bioinformatik – Methoden zur Vorhersage von RNA- und Proteinstrukturen”, Birkhäuser, chapter 8,10,11,13
- Zvelebil, M. and Baum, J. O. (2008). "Understanding Bioinformatics", Garland Science, Taylor & Francis Group, chapter 2, 11, 12 (partly)