#### Chapter 4—Amino Acids

The building blocks of proteins

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#### Amino Acid Structure

- Alpha amino acids-typical structure
  - See Figure 4.1
- Amide bond called peptide bond
  - See Figure 4.2
- Amino acids differ in R-group
  - Structures of the 20 common amino acids
  - See Figure 4.3

(LEARN structures, names, and abbreviations!!)

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## **Amino Acid Stereochemistry**

- Natural amino acids are L
  - Configuration based on glyceraldehyde
  - Fischer projection convention:

$$\begin{array}{cccc} O & & O \\ & & & & II \\ C-H & & & C-H \\ HO-C-H & & H-C-OH \\ I & & & CH_3 \\ \end{array}$$

L glyceraldehyde D glyceraldehyde (l-glyceraldehyde) (d-glyceraldehyde)

## Amino Acid Stereochemistry, con't

- Cahn, Ingold, Prelog System
  - R, S configuration (Web site explanation)
  - L glyceraldehyde is S
  - D glyceraldehyde is R
  - L alanine is S
    - See Figure Page 99
- All amino acids are S except:
  - Glycine (no chiral carbon)
  - Cysteine (why?)

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## Amino Acids are Polyprotic

- Neutral Side Chain (See Fig. 4.7)
- Acidic Side Chain (See Fig. 4.8a)
- Basic Side Chain (See Fig. 4.8b)

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## pK's of Amino Acid Groups

- See Table 1 for actual values
- · Approximate values:
  - Alpha carboxyl ~2
  - Alpha amino ~9.5
  - Side chain carboxyl ~4
  - His side chain ~6
  - Cys side chain ~8.5
  - Lys and Tyr side chain  $\sim 10$
  - Arg side chain ~12

#### Structure Effect on pK's

- · Carboxyl group
  - Alkyl carboxyl ~4.5
  - Side chain carboxy ~4
  - Alpha carboxyl ~2
  - Peptide terminal carboxy ~3
- Amino group
  - Alkyl amino ~10.5
  - Side chain amino ~10.5
  - Alpha amino ~9.5
  - Peptide terminal amino~8.

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#### **Isoionic Point**

- When  $\Sigma$  neg.charges =  $\Sigma$  pos. charges
- $pH = (pK_a + pK_b)/2$  where
  - pK<sub>a</sub> is for protonation of neutral species
  - pK<sub>b</sub> is for deprotonation of neutral species
- Illustration for neutral, acidic and basic amino acids
- Illustration for peptides

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## Determining Charge on a Peptide

- Note pK's of all functional groups
- Calculate charge when totally protonated
- Determine number of dissociable protons, and charge on each intermediate
- Determine pK's involving neutral species
- pI is average of these pK's
  - pH>pI, peptide is negatively charged
  - pH<pI, peptide is positively charged

Demonstrate with structures how electron withdrawing properties of neighboring functional groups affects the pK's.

# Determining Charge on a Peptide, con't.

- Consider the peptide **ACKRDM**
- Estimate its pI
- Estimate its charge at
  - pH = 4.0, pH = 7.0, and pH = 10.0
- How to proceed?
- First, determine the protonatable groups and their pK's

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## Functional Groups of ACKRDM

- Peptide amino terminal, pK  $\sim 8$
- Cysteine side chain, pK ~ 8.5
- Lysine side chain, pK ~ 10
- Arginine side chain, pK ~ 12
- Aspartate side chain, pK ~ 4
- Peptide carboxy terminal, pK  $\sim 3$

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## Charge of Fully Protonated ACKRDM

Total charge of fully protonated form = +3

## Charge of Fully Unprotonated ACKRDM

Total charge of fully unprotonated form = -3

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## Dissociation Steps for ACKRDM

$$+3 \xrightarrow{pK_1} +2 \xrightarrow{pK_2} +1 \xrightarrow{pK_3} 0 \xrightarrow{pK_4} -1 \xrightarrow{pK_5} -2 \xrightarrow{pK_6} -3$$

$$-3 \xrightarrow{\text{carboxyltem.}} -3 \xrightarrow{\text{aspartate amino term. cysteine lysine arginine}} -1 \xrightarrow{\text{pI}} -2 \xrightarrow{\text{pK}} -3 \xrightarrow{\text{pK}}$$

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## ACKRDM Charge at various pH's)

Group	pH = 4.0	pH = 7.0	pH = 10.0
Amino Term.	+1	~+0.9	~+0.03
Cysteine	~0	~-0.03	~-0.97
Lysine	+1	+1	~+0.5
Arginine	+1	+1	~+.99
Aspartate	~-0.5	-1	-1
Carbox. Term.	~-0.9	-1	-1
Total	~+1.6	~+0.9	~-1.5

#### Other Properties of Amino Acids

- Ultraviolet Spectra (See Figure 4.15)
- Reactions of Carboxyl Group (See Figure 4.9a)
- Reactions of Amino Group (See Figure 4.9b)
- Ninhydrin Reaction (See Figure 4.10)
- Reactions of Cysteine SH (See Figure 4.11a and 4.11b)
- <sup>13</sup>C Chemical Shifts with pH (See Figure 4.17)

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## Chromatographic Separation of Amino Acids

- Ion Exchange Chromatography
  - · Cation Exchange Resin
    - Structure (See Figure 4.18a)
    - Operation (See Figure 4.19)
  - Anion Exchange Resin (See Figure 4.18b)
- · Analysis on Ion Exchange
  - See Figure 4.21a
  - See Figure 4.21b
- HPLC Reversed Phase Chromatography of Chemical Derivative
  - See Figure 4.22