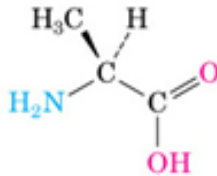


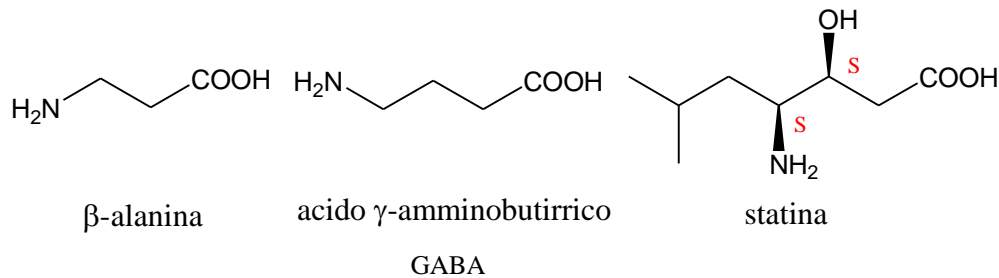
# Amminoacidi

---

Sono molecole che contengono un gruppo amminico e un'acido carbossilico. Sono stati identificati più di 700 AA diversi. Vengono classificati in funzione della posizione relativa della funzione amminica e di quella acida:  $\alpha$ ,  $\beta$ ,  $\gamma$ , etc. Hanno ruoli biologici molto diversi.



**alanina:**  $\alpha$ -amminoacido proteinogenico

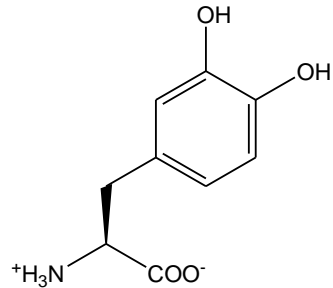


**$\beta$ -alanina:** unità strutturale presente nel CoA.

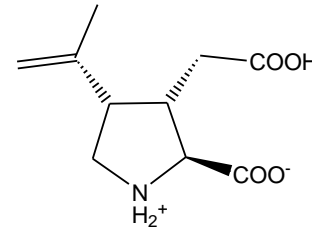
**GABA:** neurotrasmettitore ad azione inibitrice; ha azione ipertensiva

**statina:** componente di un pentapeptide che inibisce l'azione di una proteasi, la pepsina

# Aminoacidi non proteinogenici

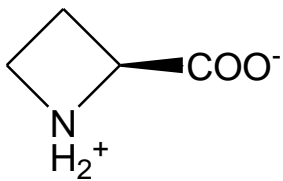


**L-Dopa**  
(anti-Parkinson's)



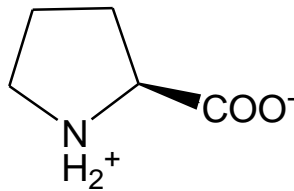
**Acido kainico**  
Eccitante (funghi)

## Aminoacidi non proteinogenici incorporati in proteine

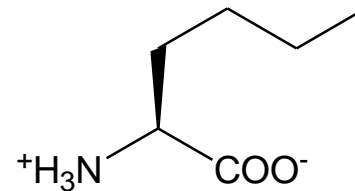


**Ac. Azetidina-2-carbossilico**

per

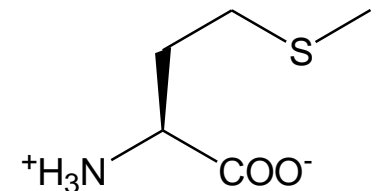


**Pro**



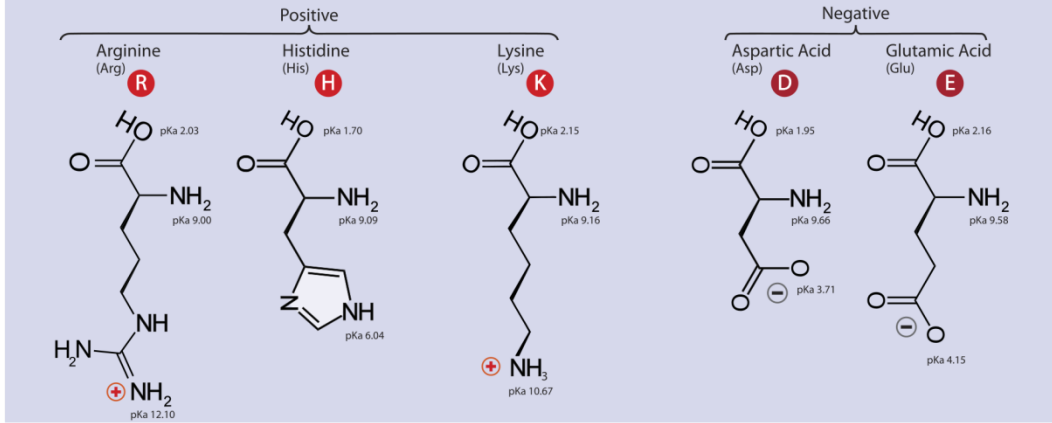
**Norleucina**

per

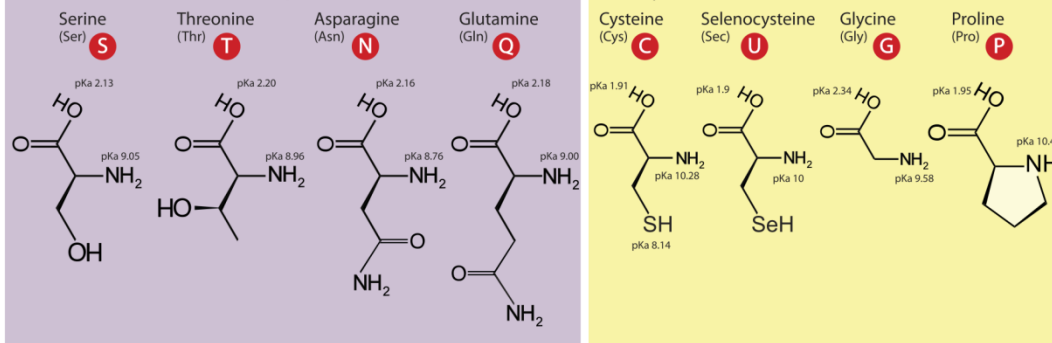


**Met**

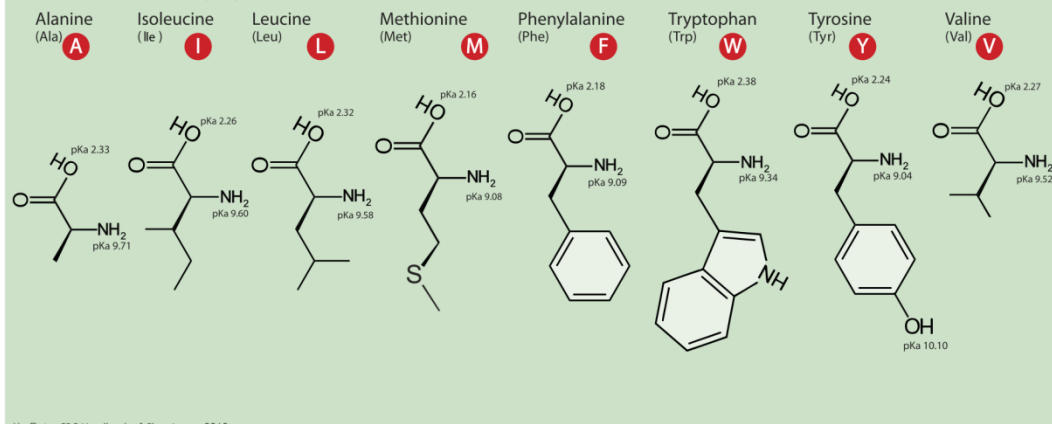
A. Amino Acids with Electrically Charged Side Chains



B. Amino Acids with Polar Uncharged Side Chains



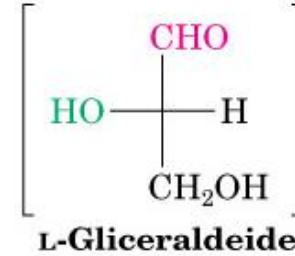
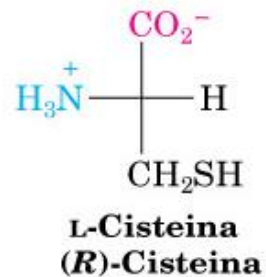
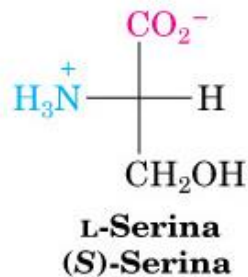
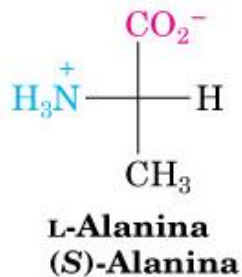
D. Amino Acids with Hydrophobic Side Chain



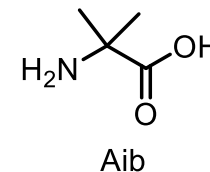
# Aminoacidi

A parte la glicina tutti gli amminoacidi proteinogenici sono chirali e, negli organismi superiori, le proteine sono formate solo da amminoacidi della serie L.

Negli organismi inferiori (batteri) si trovano anche amminoacidi della serie D e amminoacidi non chirali come l'Aib (acido  $\alpha$ -amminoisobutirrico)



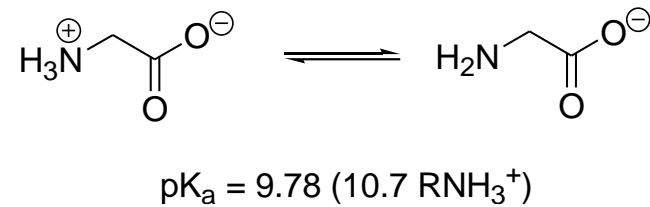
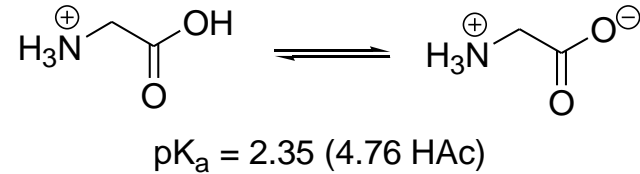
(L)A-(L)A-(L)A-(L)A-(L)A-(L)A  
(D)A-(L)A-(L)A-(L)A-(L)A-(L)A  
(L)A-(D)A-(L)A-(L)A-(L)A-(L)A  
etc, etc ....



# Aminoacidi

TABELLA 27.2 VALORI DEI  $pK_a$  PER I GRUPPI IONIZZABILI DEGLI AMMINOACIDI

Amminoacido	$pK_a$ $\alpha$ -CO <sub>2</sub> H	$pK_a$ $\alpha$ -NH <sub>3</sub> <sup>+</sup>	$pK_a$ catena laterale	punto isoelettrico (pI)
alanina	2.35	9.87	—	6.11
arginina	2.01	9.04	12.48	10.76
asparagina	2.02	8.80	—	5.41
acido aspartico	2.10	9.82	3.86	2.98
cisteina	2.05	10.25	8.00	5.02
acido glutammico	2.10	9.47	4.07	3.08
glutammina	2.17	9.13	—	5.65
glicina	2.35	9.78	—	6.06
istidina	1.77	9.18	6.10	7.64
isoleucina	2.32	9.76	—	6.04
leucina	2.33	9.74	—	6.04
lisina	2.18	8.95	10.53	9.74
metionina	2.28	9.21	—	5.74
fenilalanina	2.58	9.24	—	5.91
prolina	2.00	10.60	—	6.30
serina	2.21	9.15	—	5.68
treonina	2.09	9.10	—	5.60
triptofano	2.38	9.39	—	5.88
tirosina	2.20	9.11	10.07	5.63
valina	2.29	9.72	—	6.00



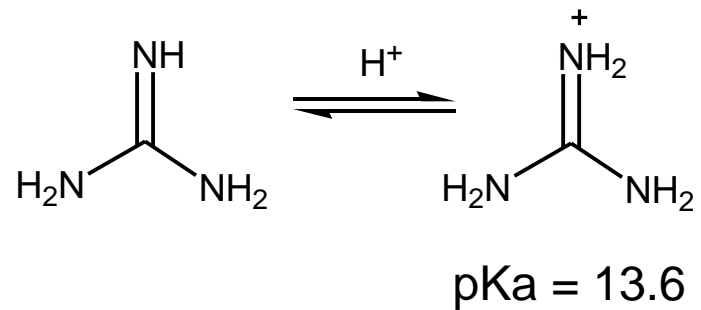
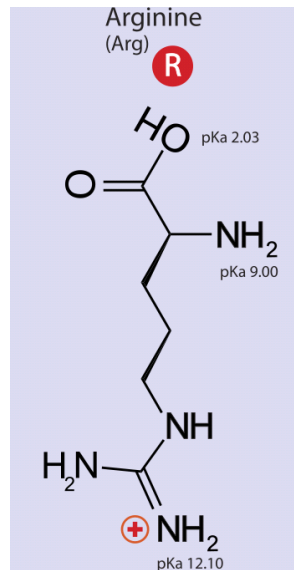
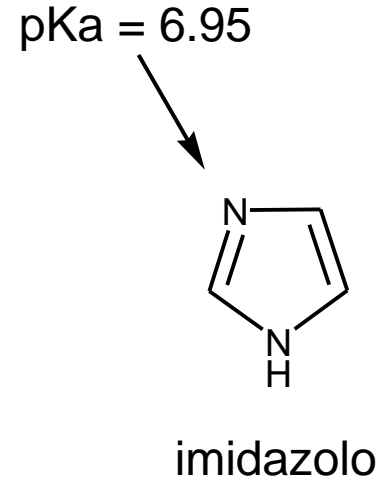
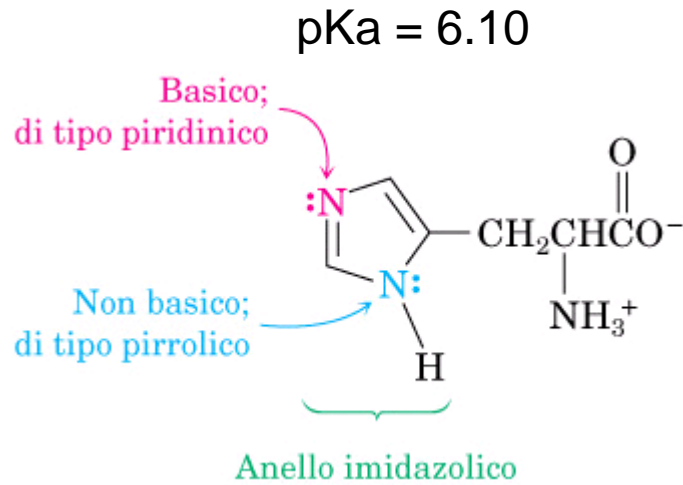
## **Punto Isoelettrico (pI)**

E' il valore di pH al quale la carica netta di una certa specie elementare è zero.

Il pI è la media dei valori di  $pK_a$  dei gruppi carbossilico e ammonio.

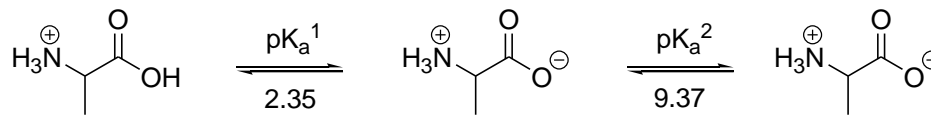
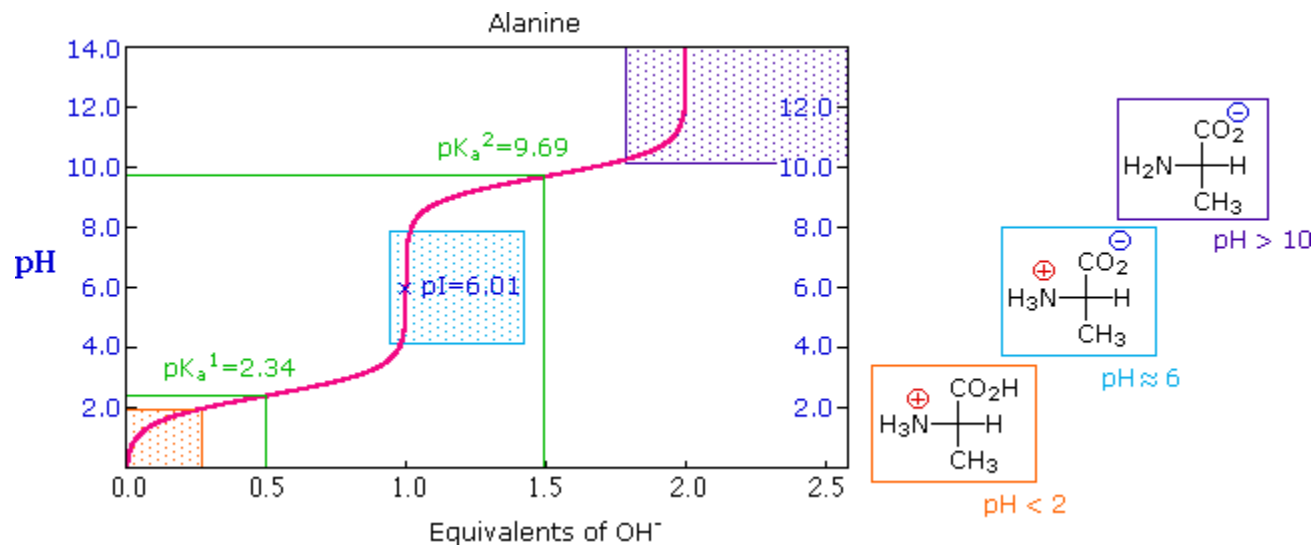
$$(9.78+2.35)/2 = 6.06$$

# Aminoacidi: istidina e arginina



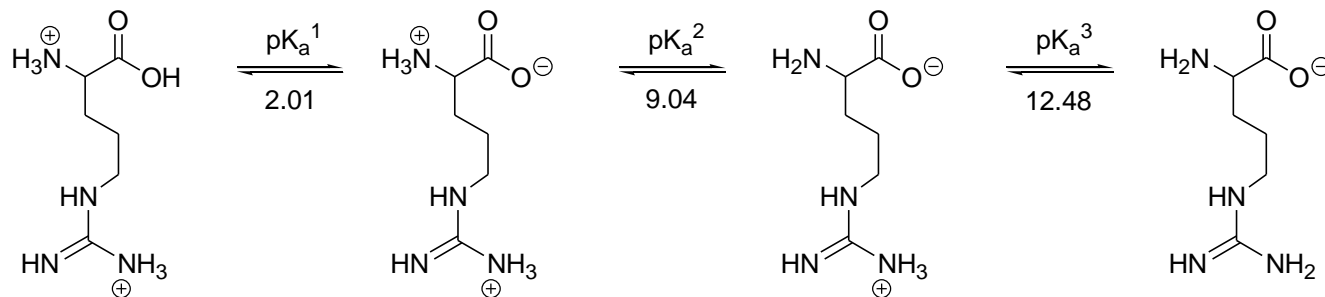
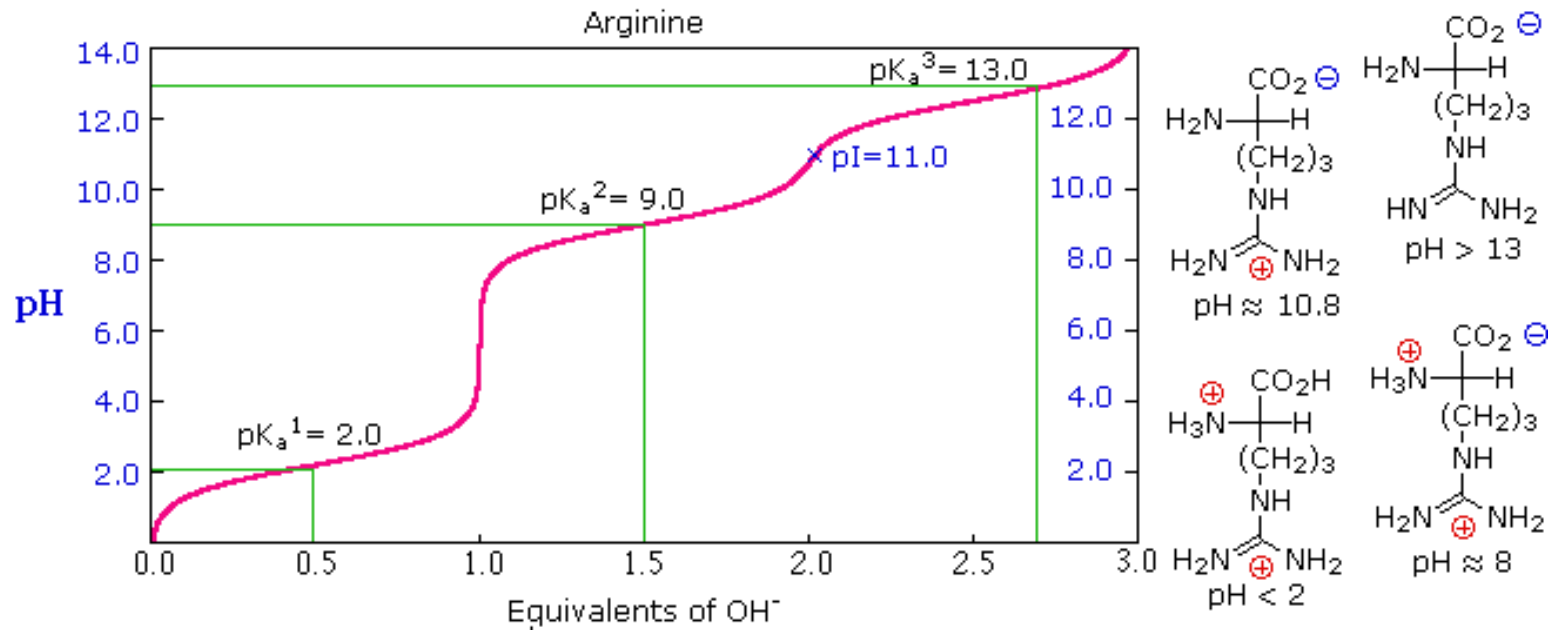
TEA pKa = 10.75

# Titolazioni di amminoacidi: ALANINA



$\text{pI} = 6.11$

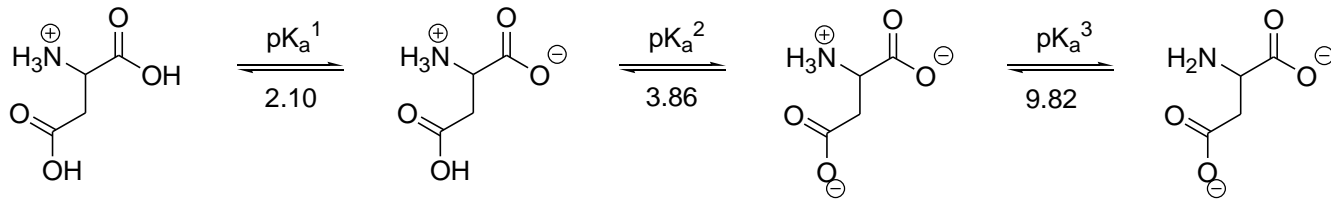
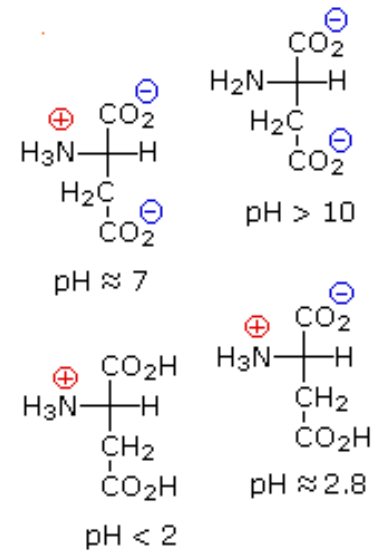
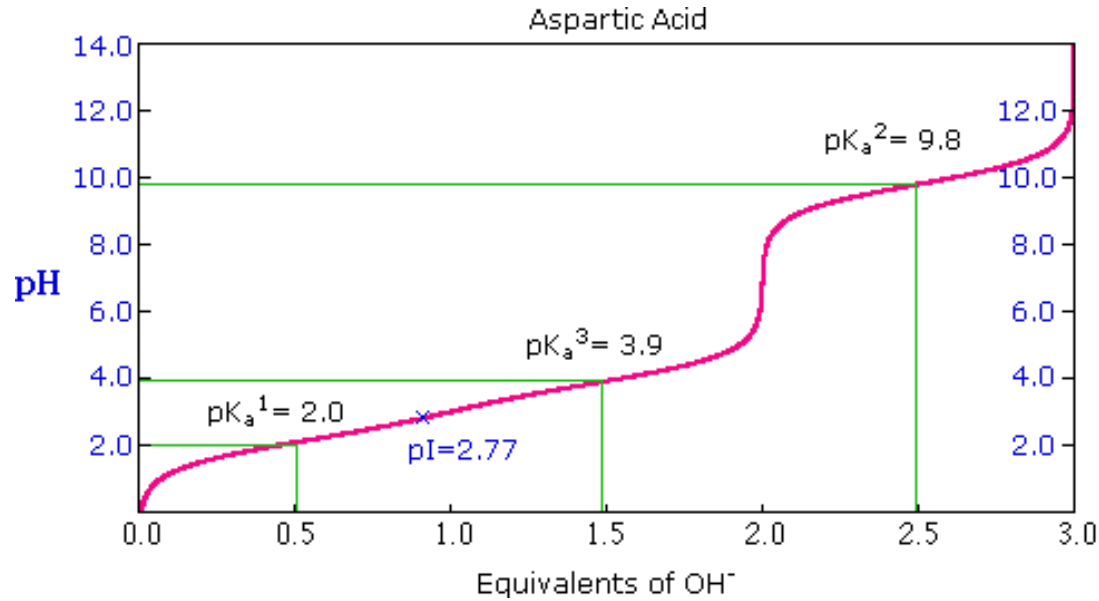
# Titolazioni di amminoacidi: Arginina



$pI = 10.7$

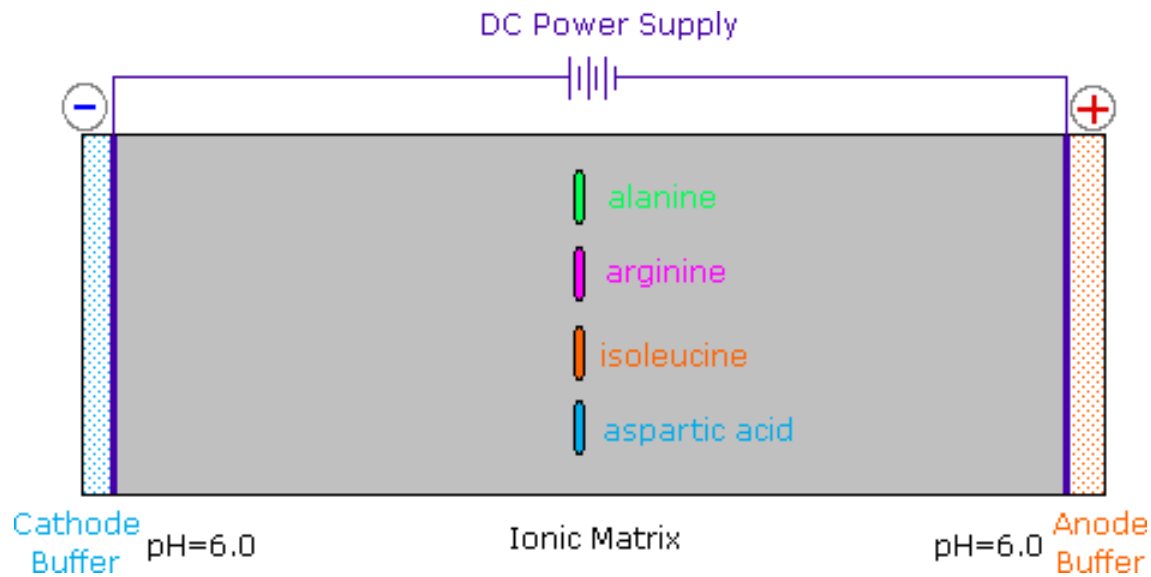


# Titolazioni di amminoacidi: Acido Aspartico

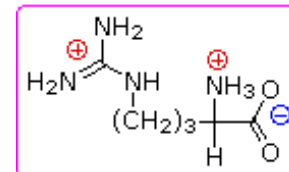


$pI = 2.98$

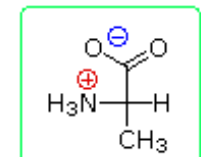
# Elettroforesi



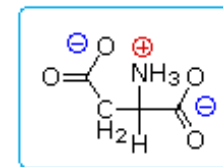
Predominant Species at pH=6.0



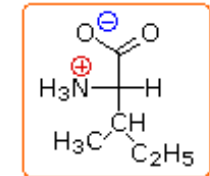
arginine  $pI=10.77$



alanine  $pI=6.01$



aspartic acid  $pI=2.80$

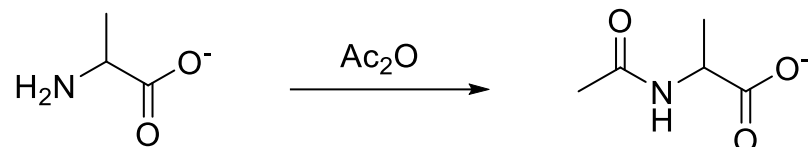


isoleucine  $pI=6.02$

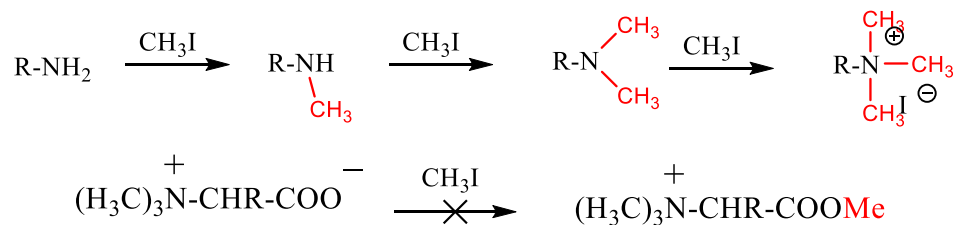
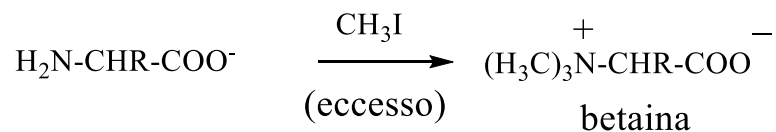
# Amminoacidi: reazioni

## Reazioni del gruppo amminico:

### acilazione



### alchilazione

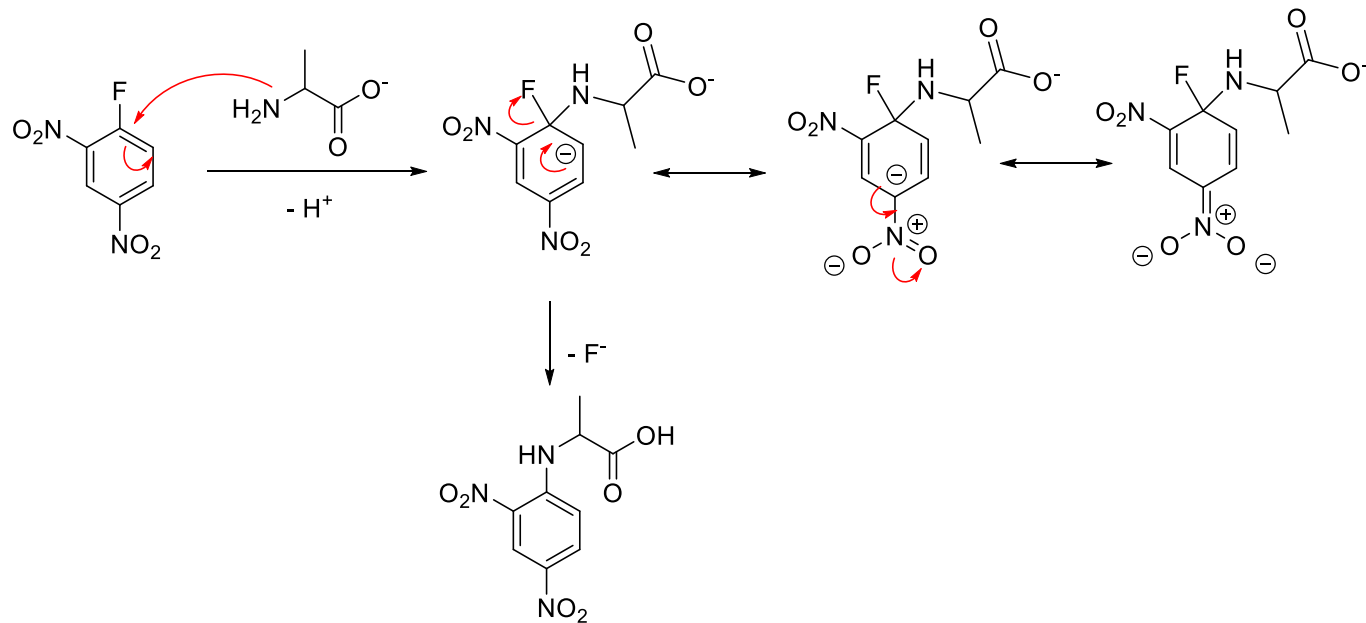
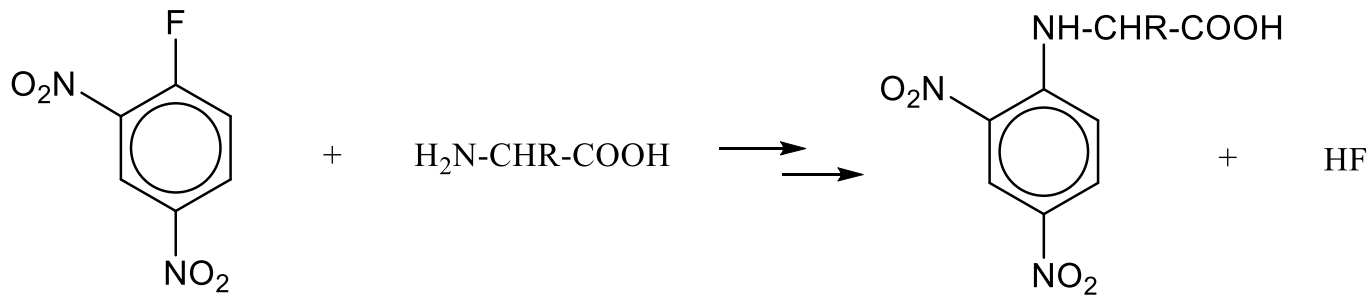


# Amminoacidi: reazioni

## Reazioni del gruppo amminico:

### alchilazione

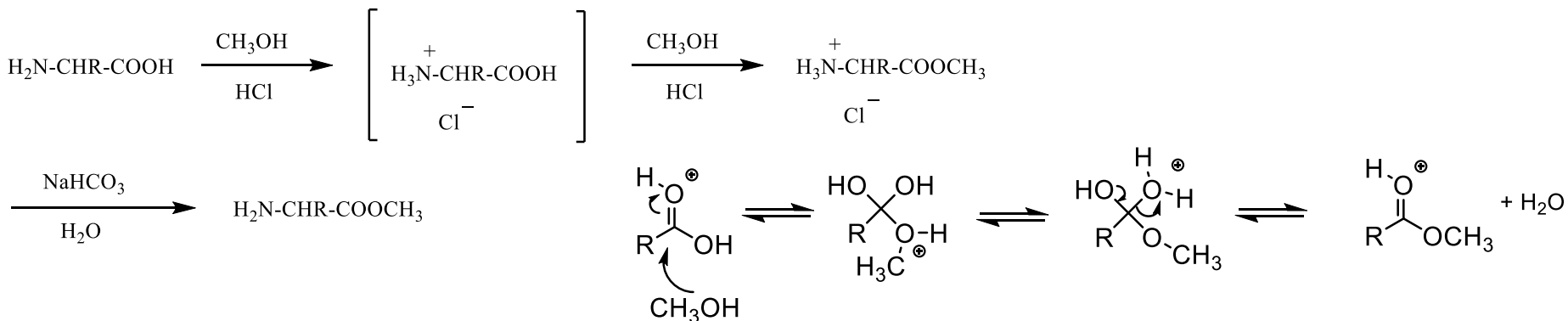
#### - reazione di Sanger



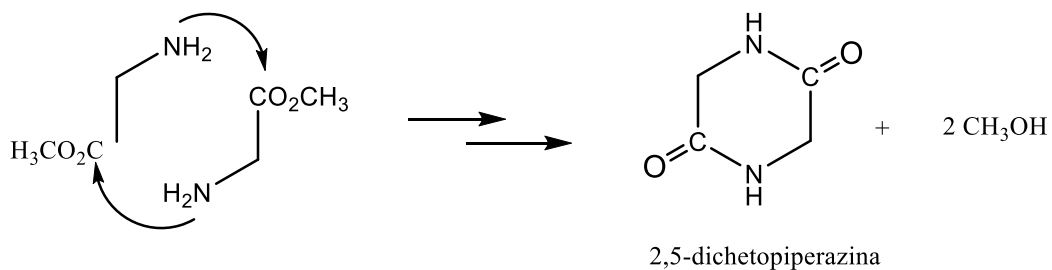
# Amminoacidi: reazioni

## Reazioni del gruppo carbossilico:

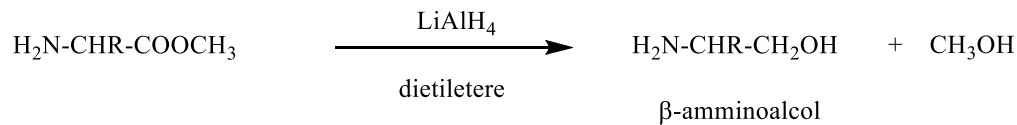
### Formazione di esteri:



NB: gli esteri degli  $\alpha$ -amminoacidi sono instabili. Più stabili i loro cloridrati.

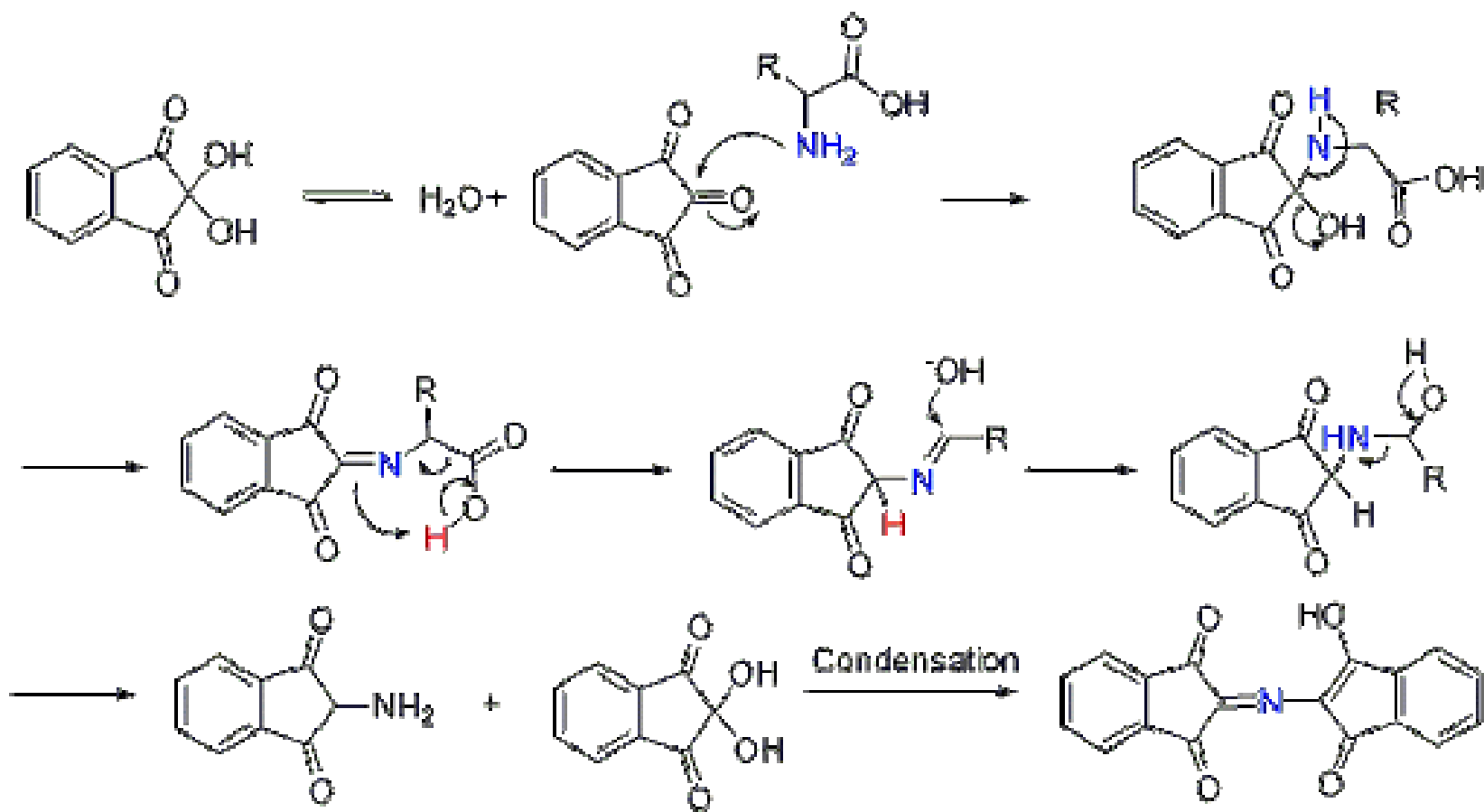


### Riduzione del gruppo estereo:



# Amminoacidi: reazioni

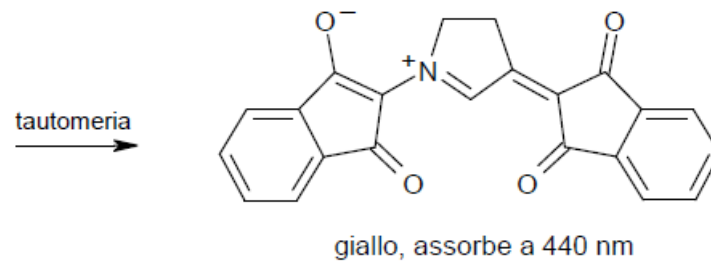
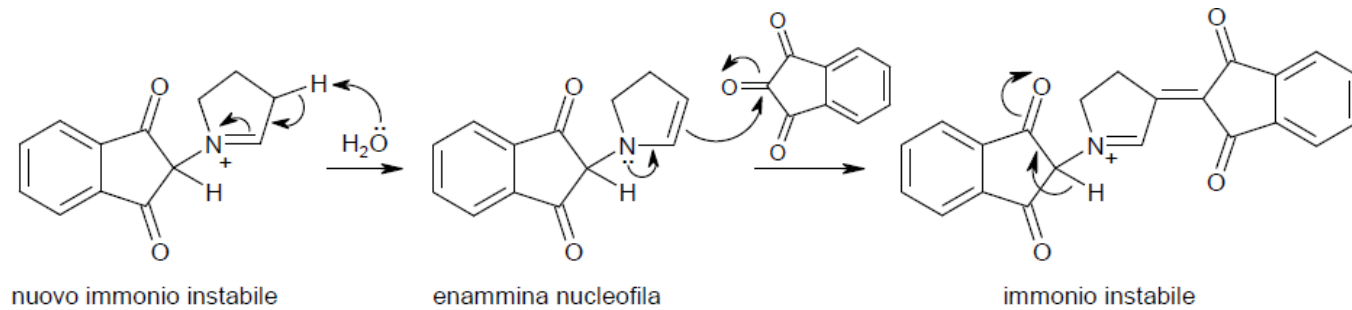
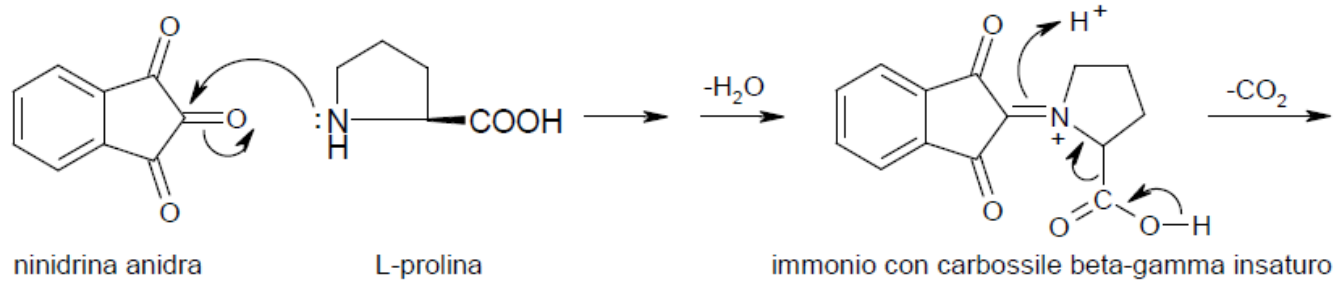
## Reazione con la ninidrina



blu magenta assorbe a 570 nm

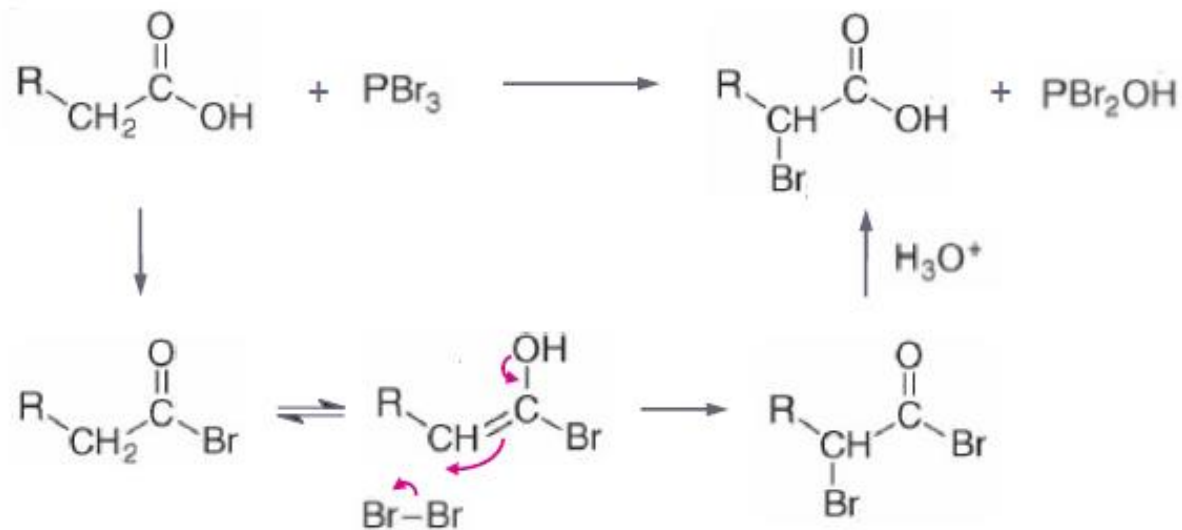
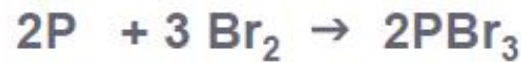
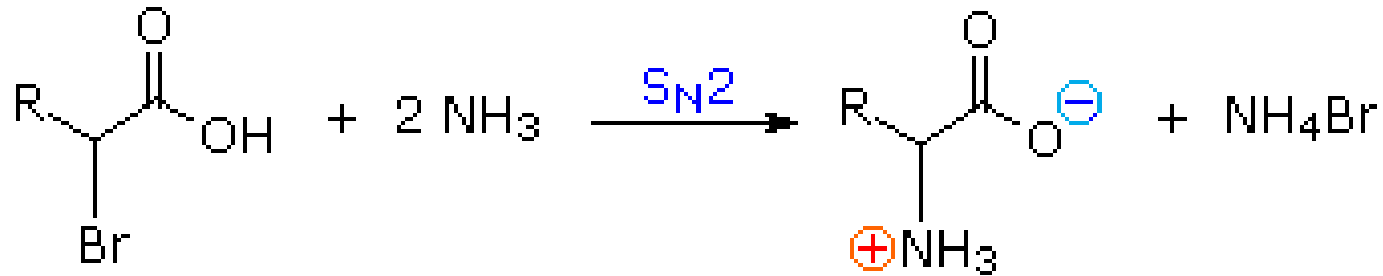
# Amminoacidi: reazioni

## Reazione con la ninidrina



# Amminoacidi: sintesi

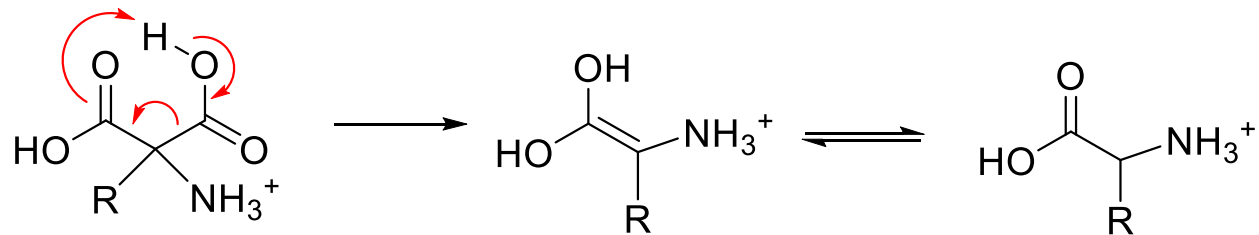
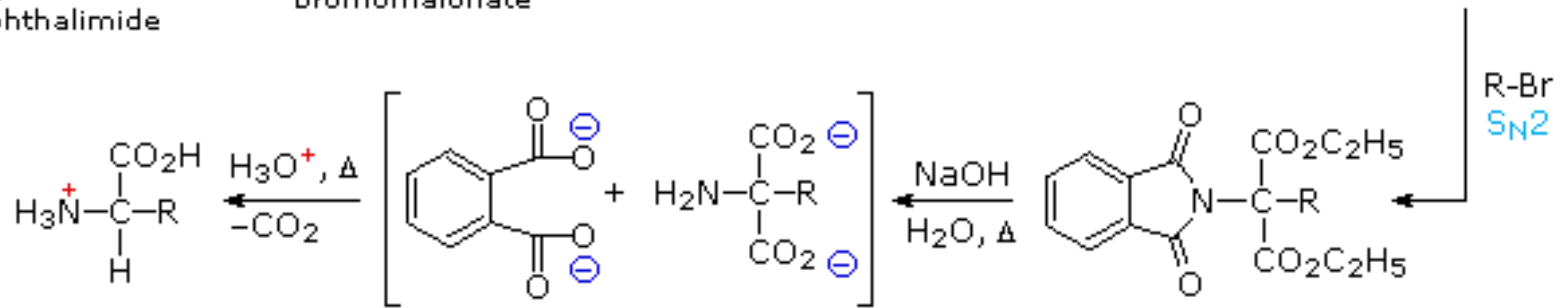
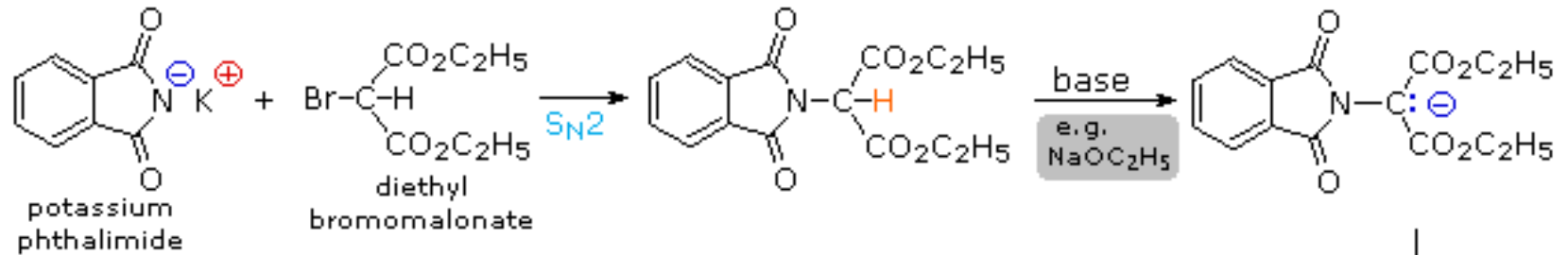
## Ammonolisi di acidi $\alpha$ -bromocarbossilici





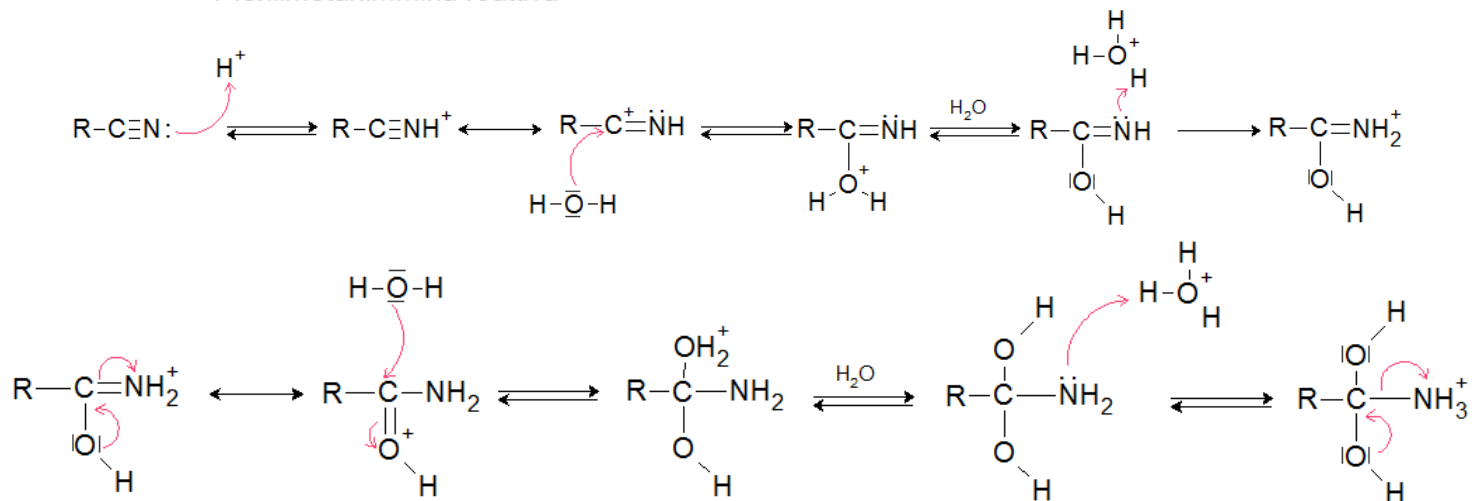
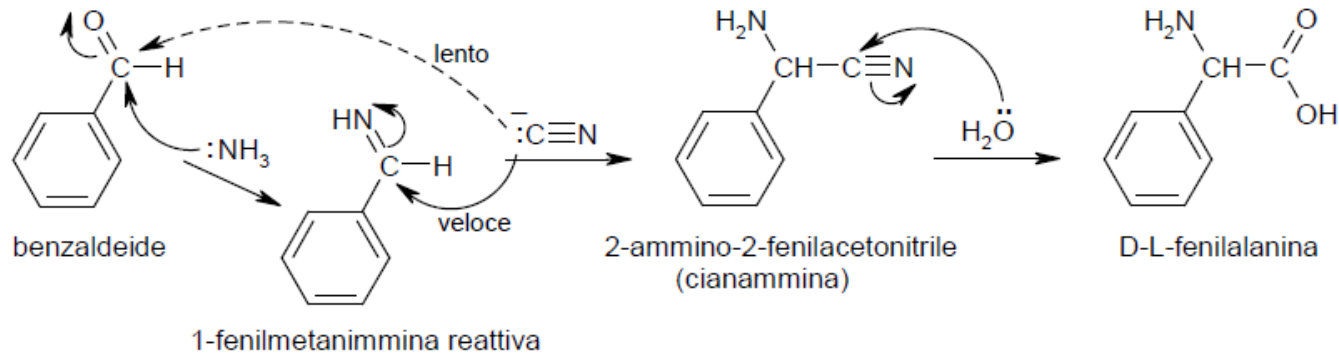
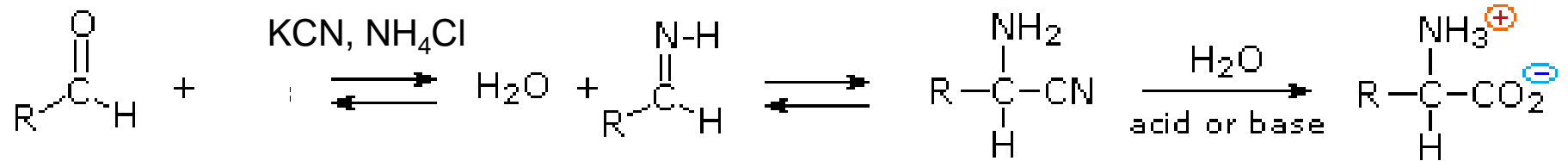
# Amminoacidi: sintesi

## Sintesi di Gabriel



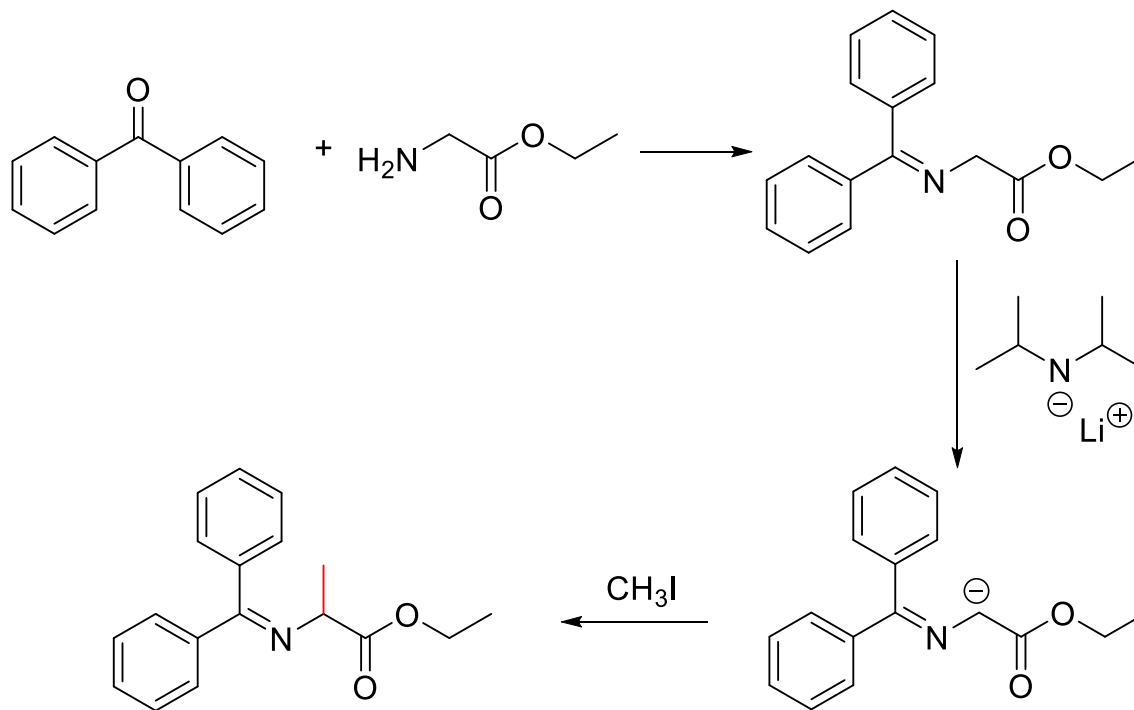
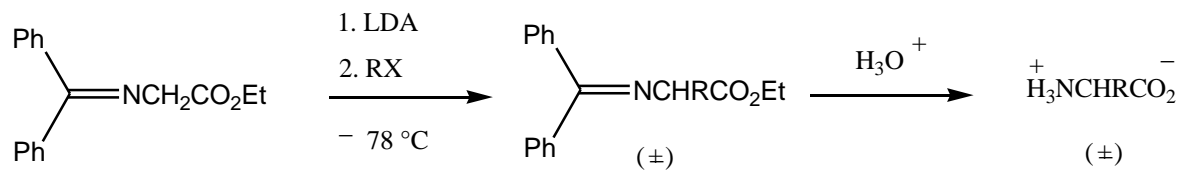
# Amminoacidi: sintesi

## Sintesi di Strecker



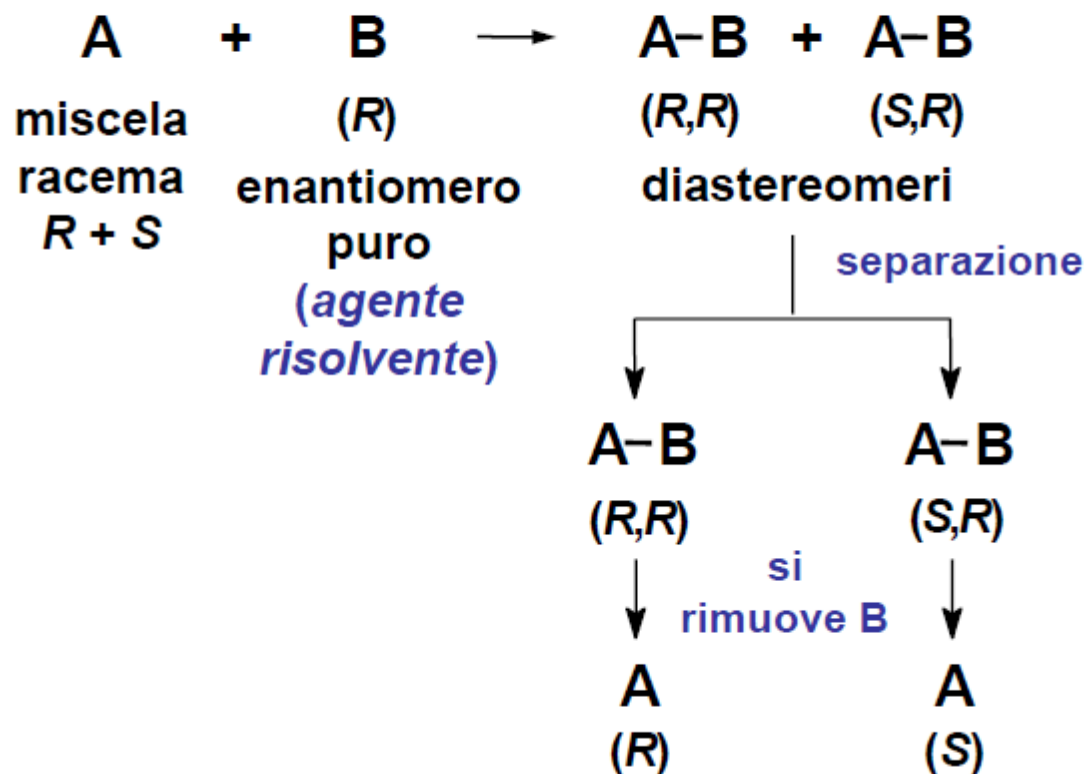
# Amminoacidi: sintesi

## Alchilazione della glicina N-funzionalizzata

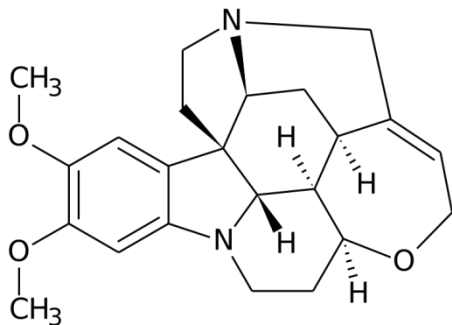
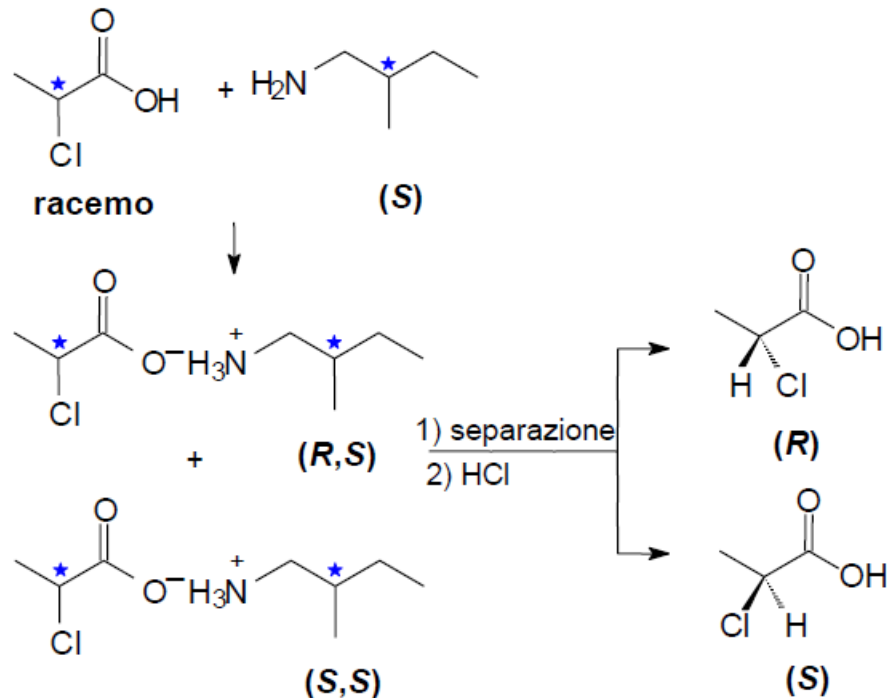


# Risoluzione di Enantiomeri

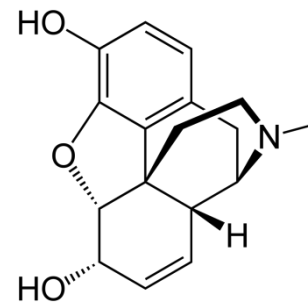
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# Risoluzione di Enantiomeri

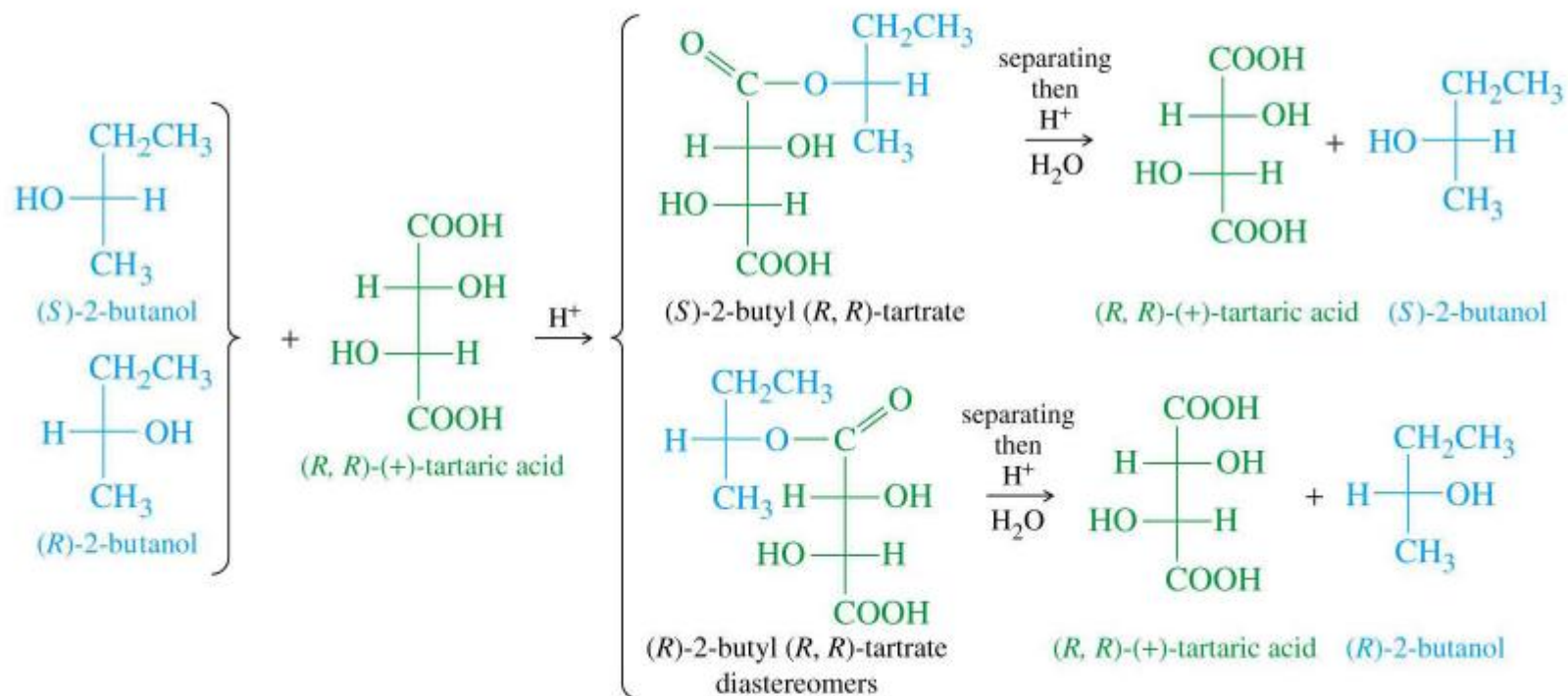


Brucina (2,3-dimetossistricnina)

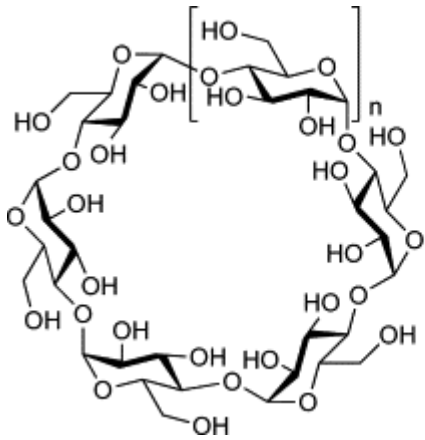


morfina

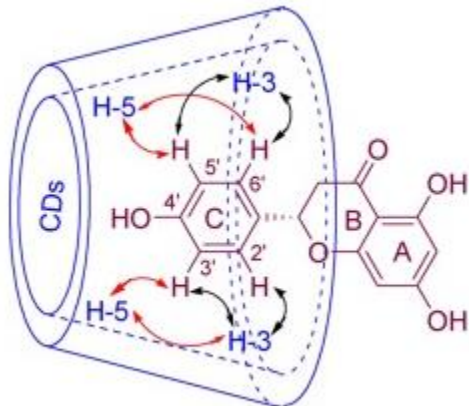
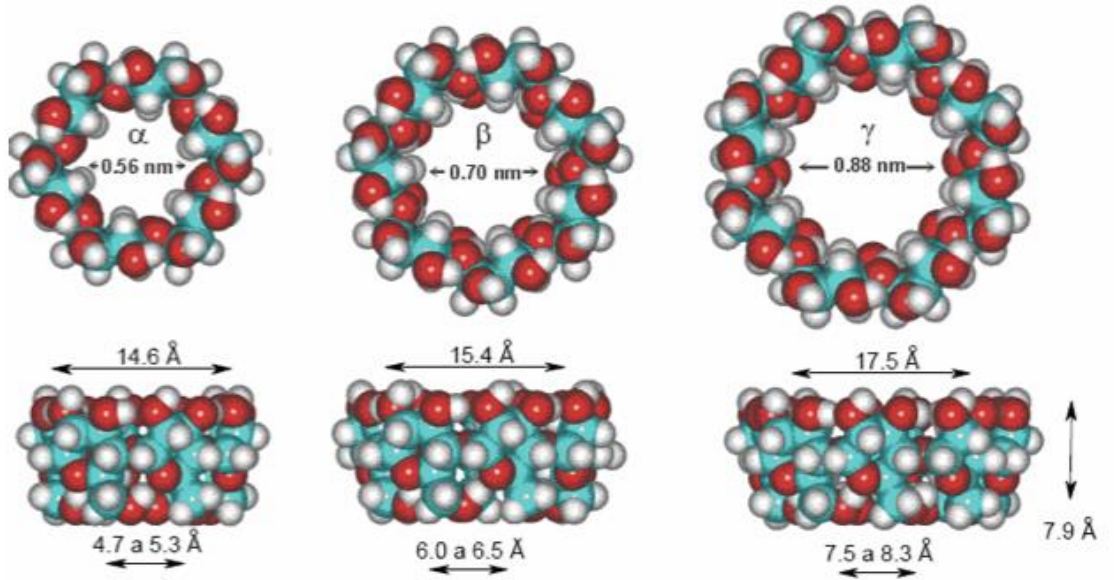
# Risoluzione di Enantiomeri



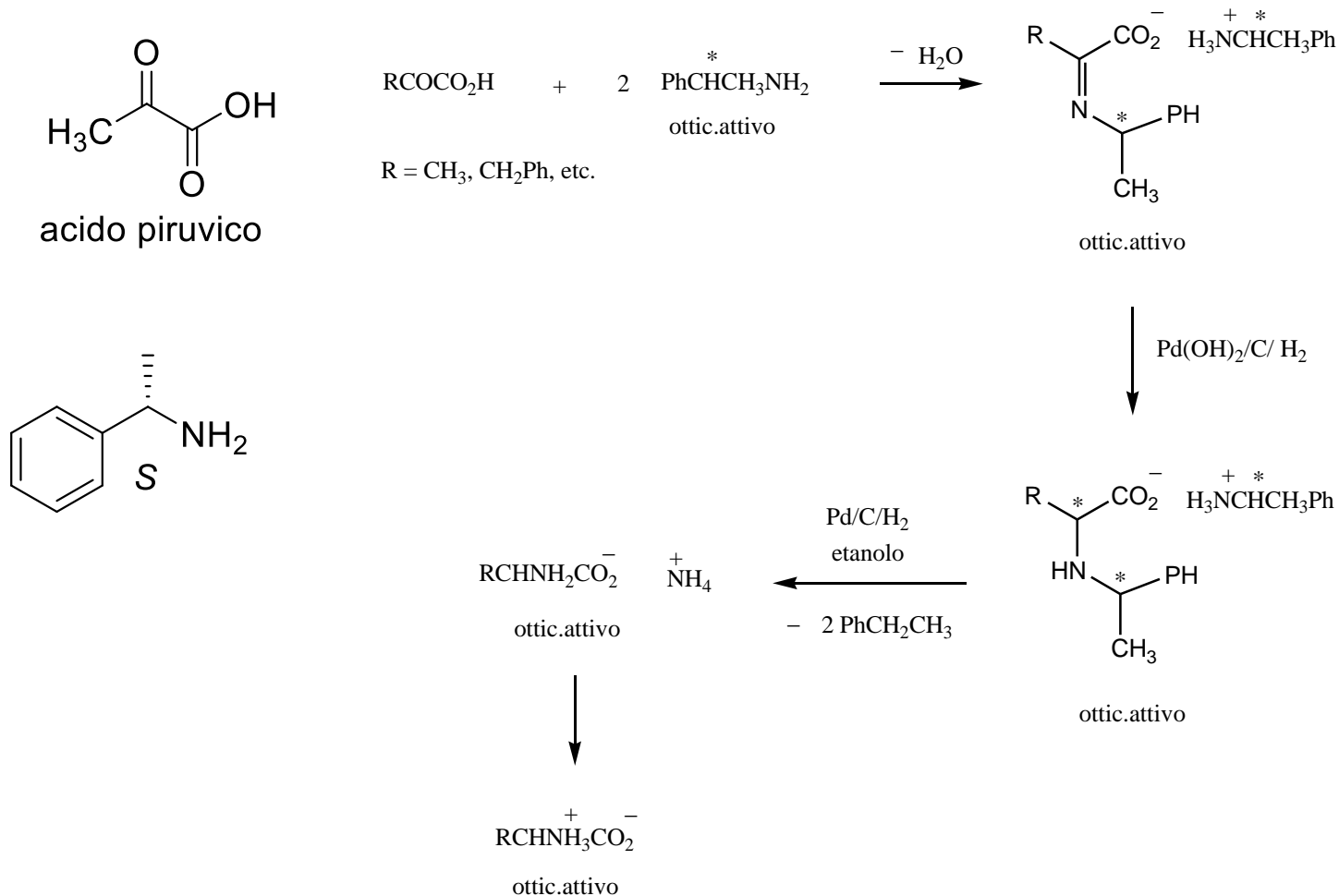
# Ciclodestrina



n = 1,  $\alpha$ -cyclodextrin  
n = 2,  $\beta$ -cyclodextrin  
n = 3,  $\gamma$ -cyclodextrin



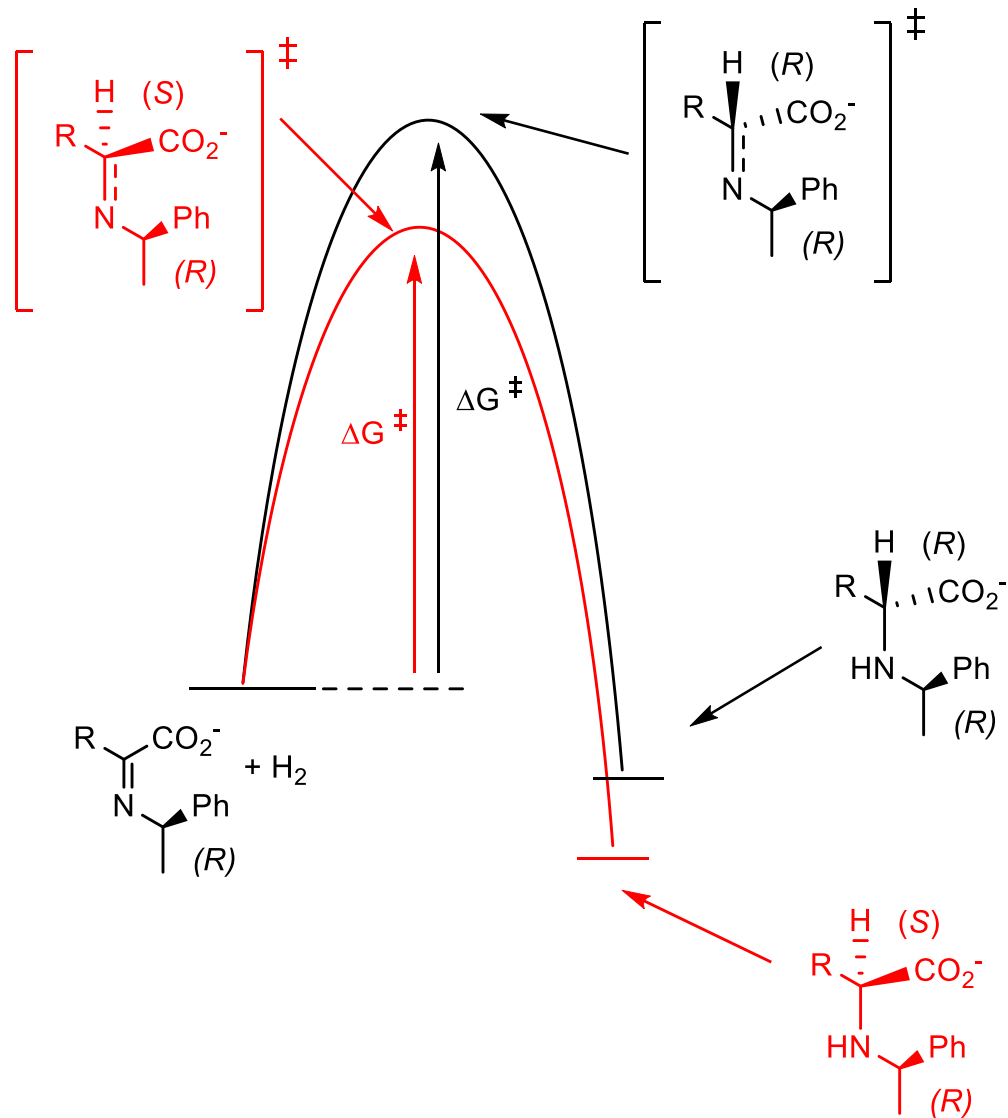
# Amminoacidi: sintesi enantioselettiva



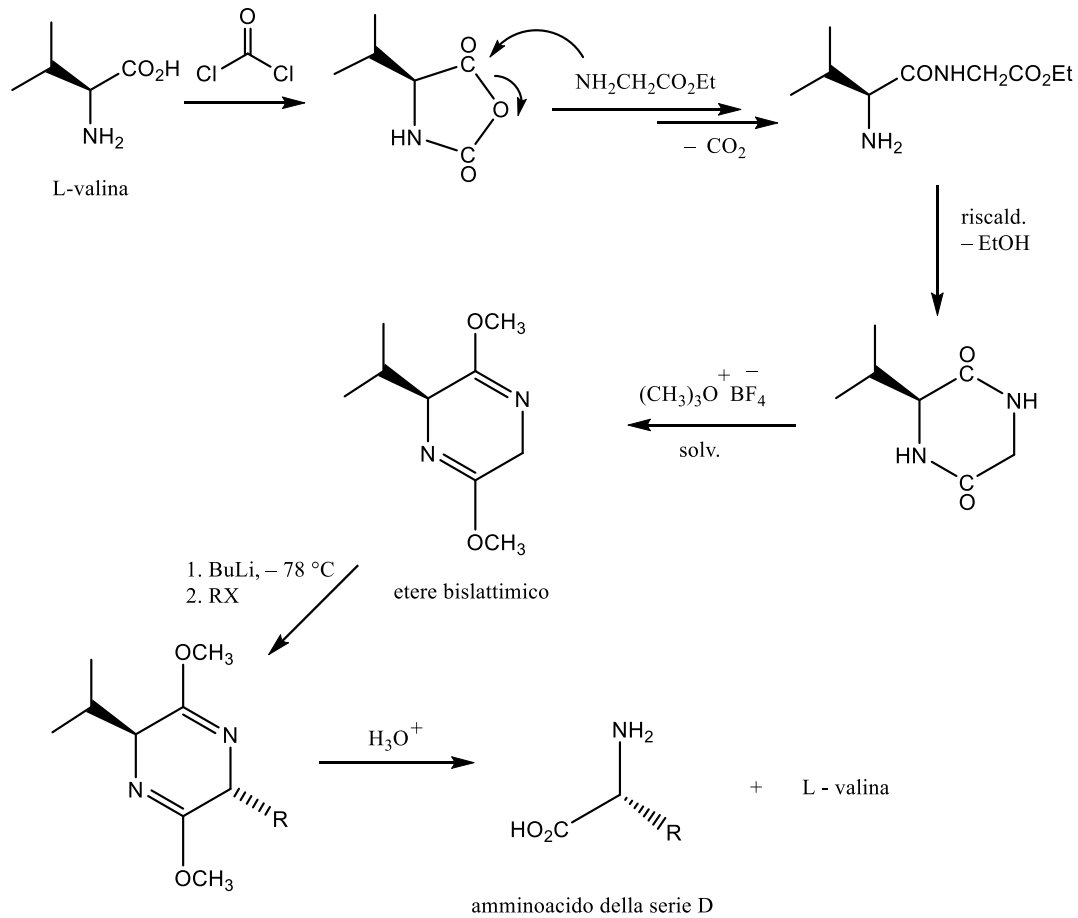
Ad es. utilizzando la (S)-(-)-1-fenetilammina, dopo l'idrogenazione del doppio legame imminico, si ottengono due diastereomeri salini in proporzioni diverse che possono essere separati per cristallizzazione frazionata, sottoposti ad idrogenolisi per dare l'amminoacido che sarà otticamente attivo.



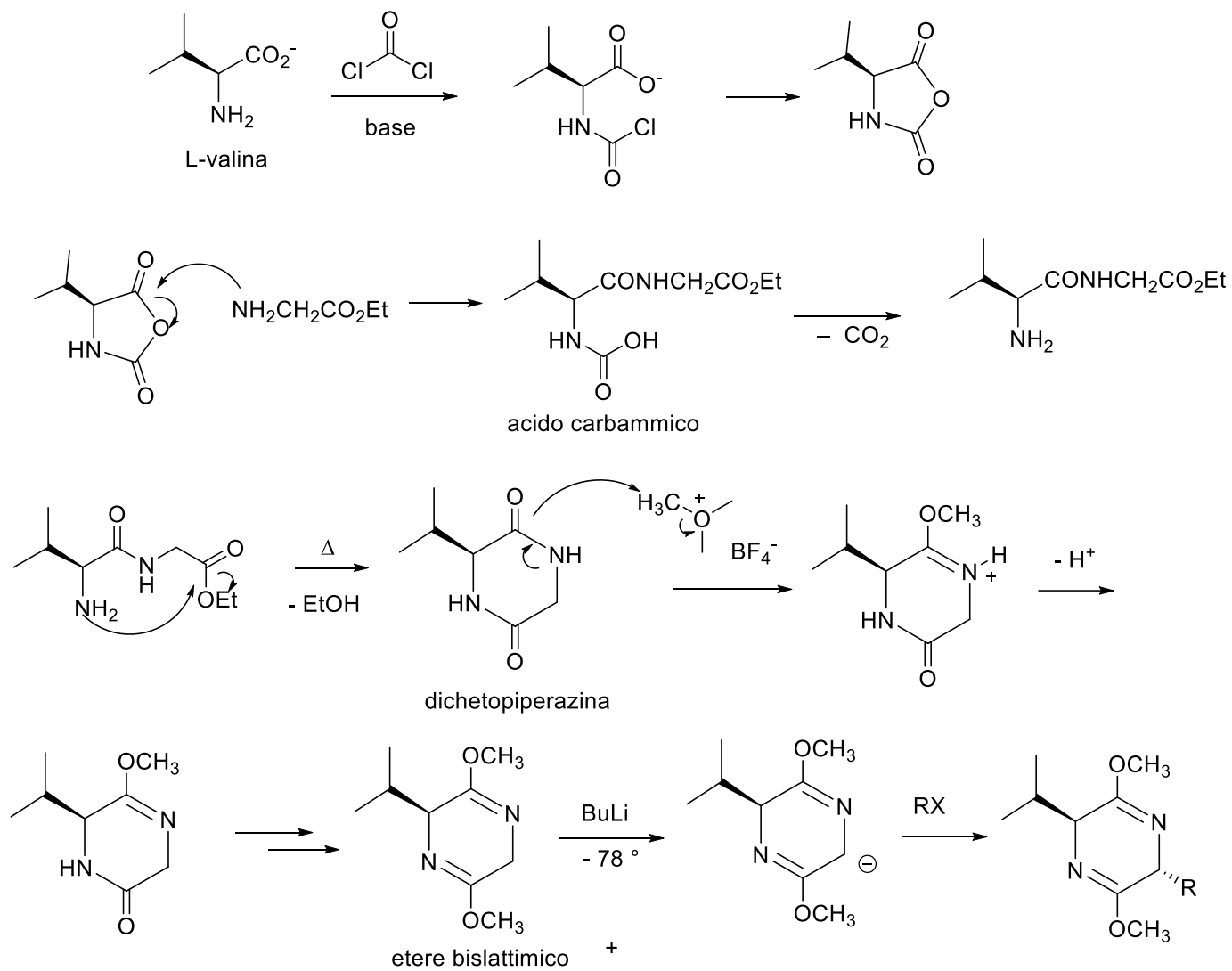
# Amminoacidi: sintesi enantioselettiva



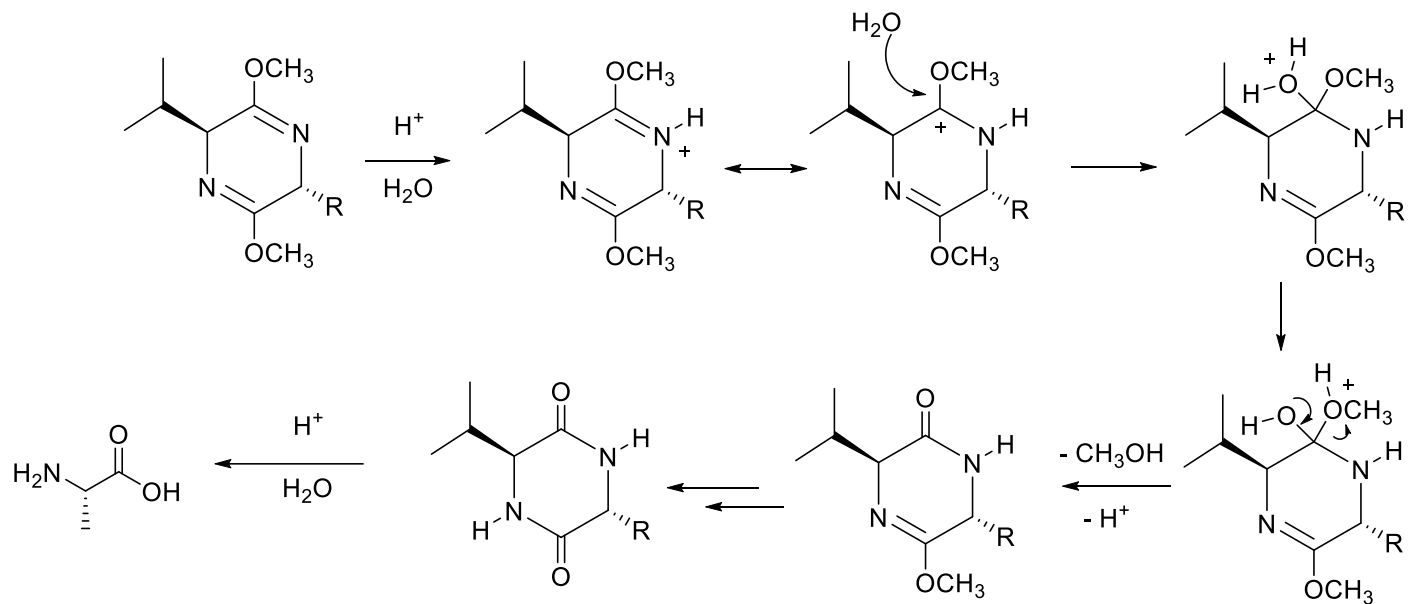
# Amminoacidi: sintesi enantioselettiva



# Amminoacidi: sintesi enantioselettiva

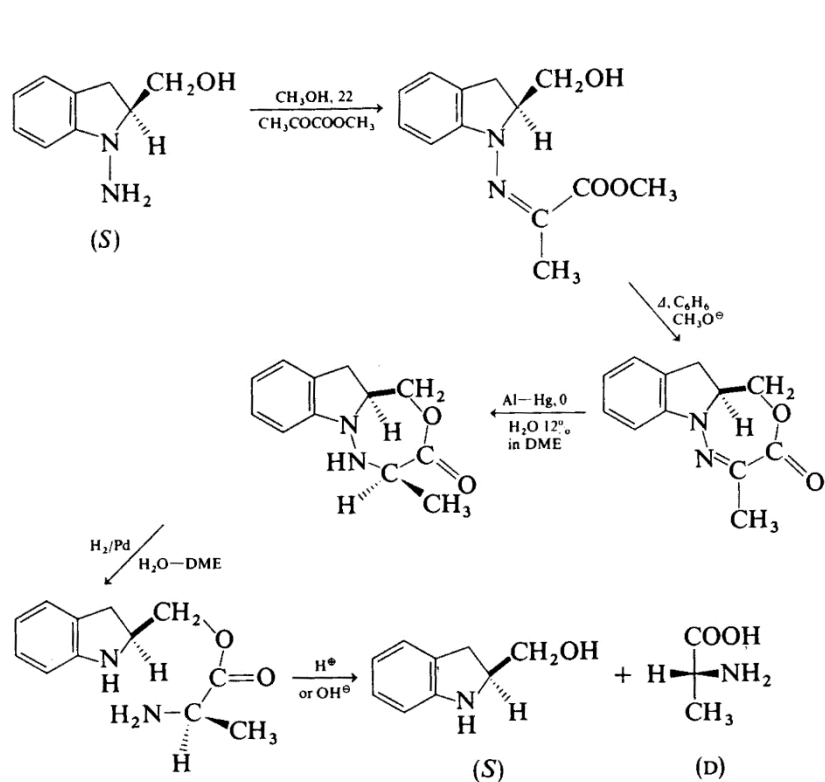


# Amminoacidi: sintesi enantioselettiva

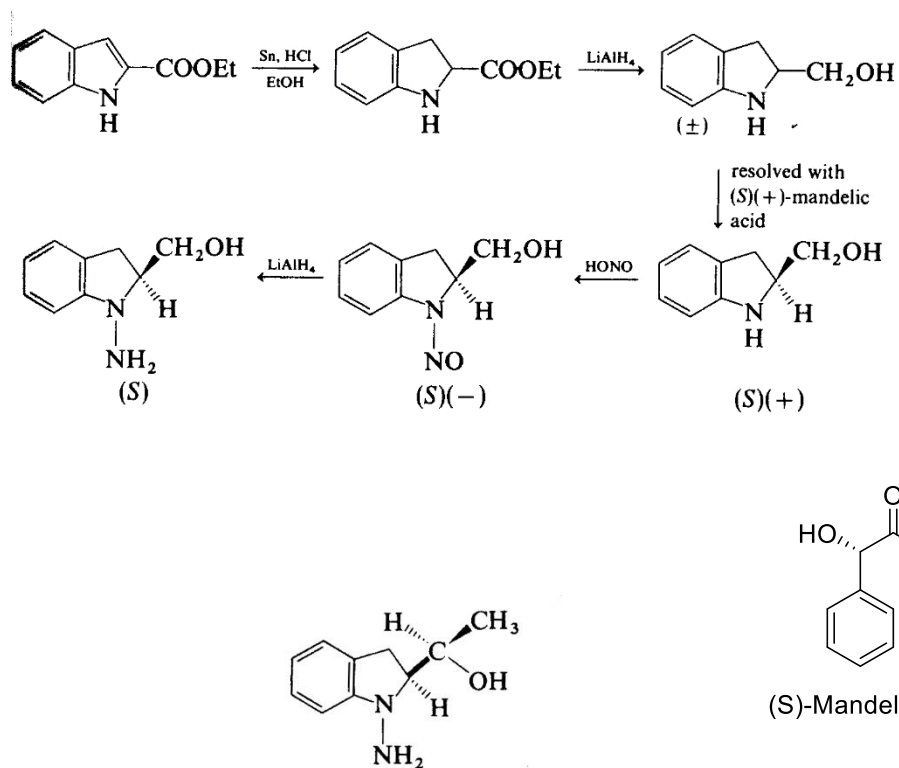


# Amminoacidi: sintesi enantioselettiva

## Sintesi di E.J. Corey



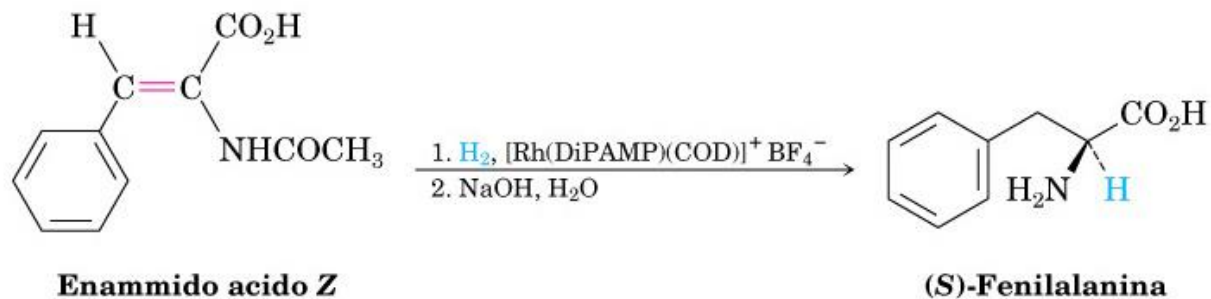
e.e. 80 %



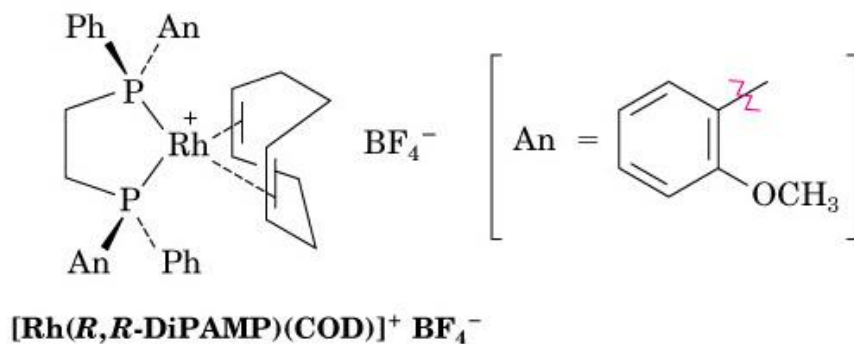
e.e. 96 %

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## Sintesi di W. S. Knowles

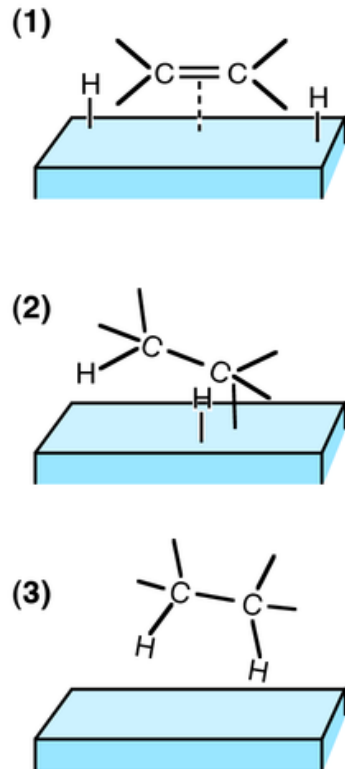


**e.e. 99 %**

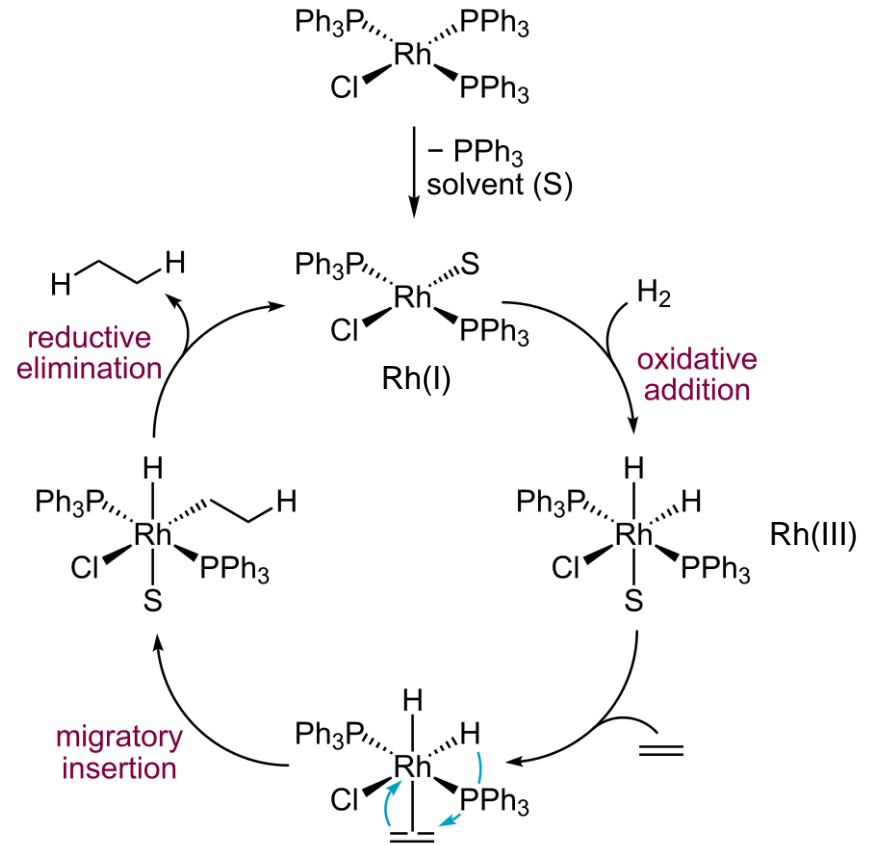


# Idrogenazione catalitica

## Catalisi eterogenea



## Catalisi omogenea

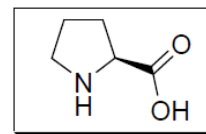
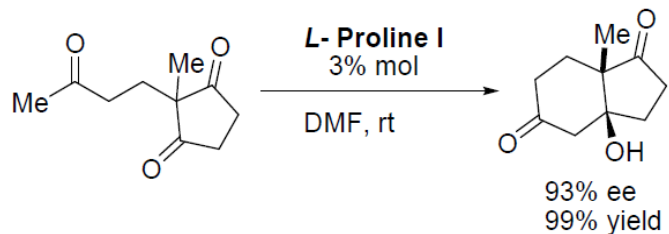


Catalizzatore di Wilkinson

**Addizione ossidativa:** un substrato viene addizionato ad un complesso metallico con il metallo che aumenta lo stato di ossidazione e il numero di coordinazione di due unità. **L'eliminazione riduttiva** è la reazione opposta.

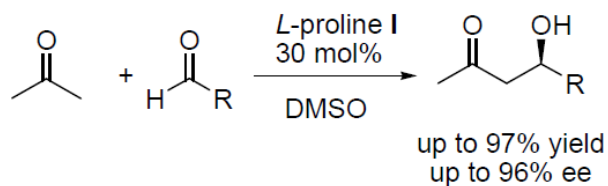
# Organocatalisi

## Ammino catalisi

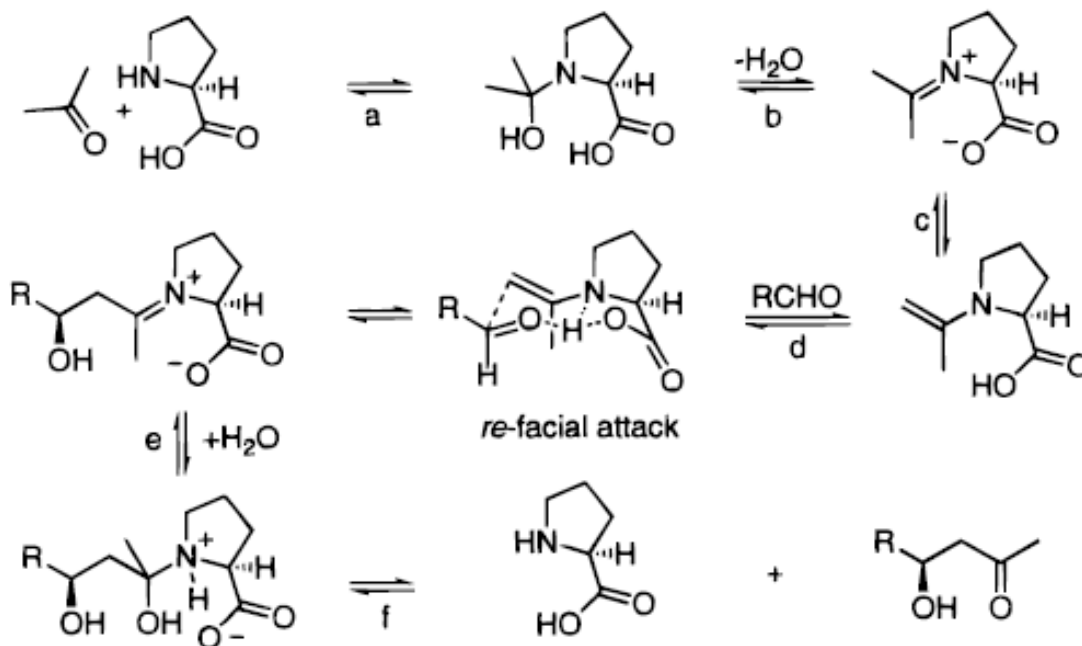


**L-Proline**

Hajos Parrish Wiechert, Eder, and Sauer, **1971**



B.List, R. A. Lerner, F. C. Barbas III *J. Am. Chem. Soc.* **2000**, *122*, 2395.





# Organocatalisi

## Ammino catalisi

