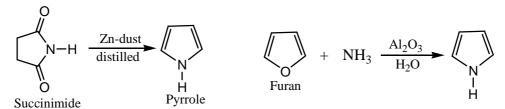
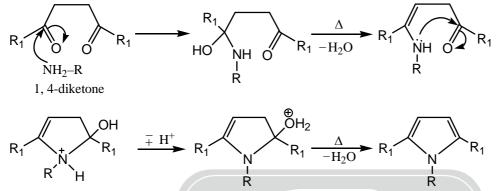
## (III) From Succinamide

## (IV) From Furan



## (V) Paal-Knorr synthesis:

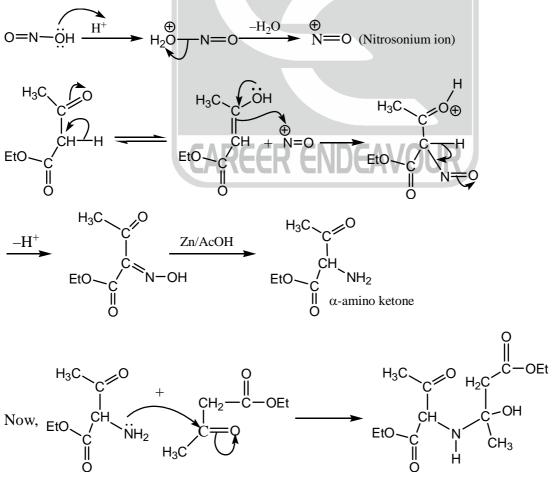
By treating 1, 4-diketone with ammonia, primary amine or hydrazine etc.



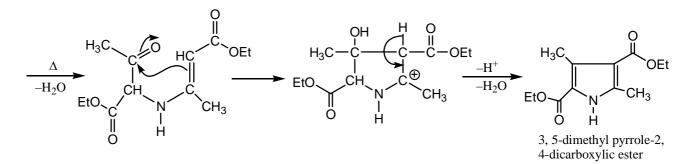
## (VI) Knorr Pyrrole synthesis:

It involves the condensation between an  $\alpha$  – amino ketone and a  $\beta$  – diketone or  $\beta$  – ketoester which produces derivative of pyrrole.

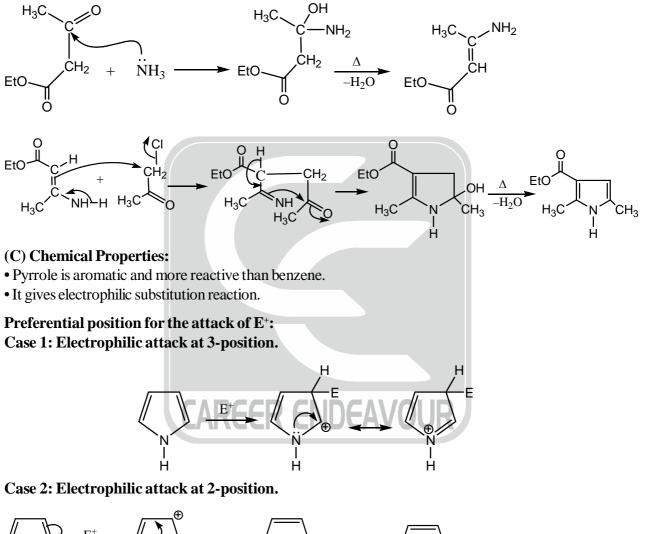
Mechanism:

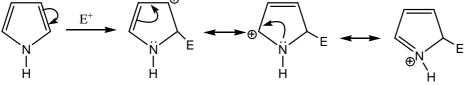






(7) Hantzsch synthesis: Condensation between chloroacetone, a  $\beta$  – ketoester and primary amine or amonia.





Since intermediate formed after the attack of  $E^+$  at 2-position is more stable (due to more resonating structure) than that of intermediate formed after electrophilic attack at 3-position so, electrophilic substitution are favorably occured at 2-position rather than three. If the 2-position is occupied then next substitution will be at 5-carbon if both position 2 and 5 are occupied then substitution can be possible at 3-position.

(1) Reaction with  $Br_2$ : Reaction with bromine requires no Lewis acid and leads to substitution at all four free positions.

