

ANTI-TUBERCULAR AGENTS

CLASSIFICATION :

BASED ON THEIR DOSAGE
REGIMEN

FIRST LINE DRUGS

ISONIAZID, STREPTOMYCIN, RIFAMPICIN,
ETHAMBUTOL, PYRAZINAMIDE

SECOND LINE DRUGS

ETHIONAMIDE

PAS P. AMINO SALICYLIC ACID

OFLOXACIN - Fluoro Quinolone

CIPROFLOXACIN - "

CYCLOSERINE - Amino acid derivative

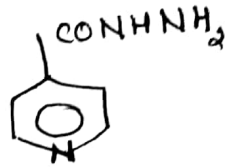
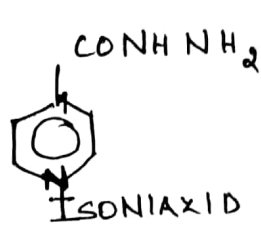
AMIKACIN

KANAMYCIN

VIOMYCIN

CAPREOMYCIN

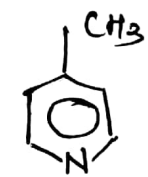
(2)



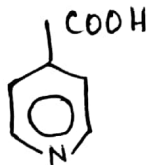
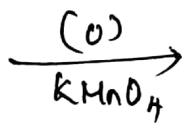
Pyridine derivative

M.O.A : Inhibits the synthesis of Mycolic acid essential for the cell wall synthesis of Bacteria

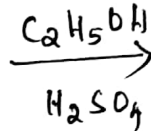
Syn



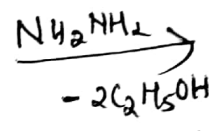
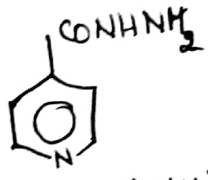
2-picolone



Isonicotinic acid



Ester

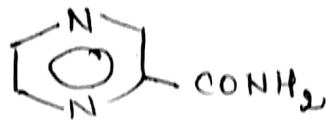
(Rxn of
O=C compound)

(INH)

Uses : Both Bacteriostatic and Bactericidal depending On the concentration.

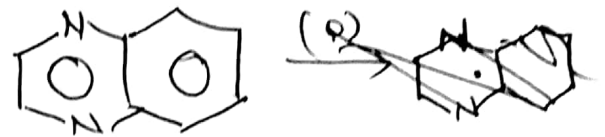
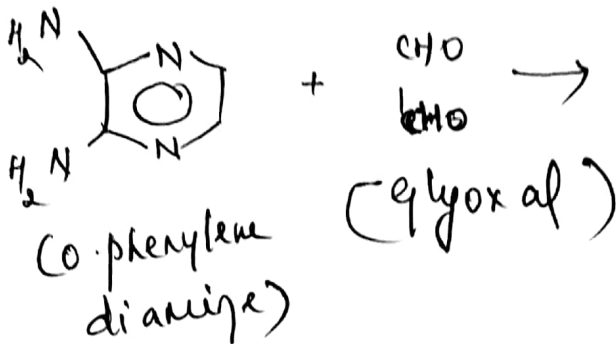
Pyrazinamide

(3)



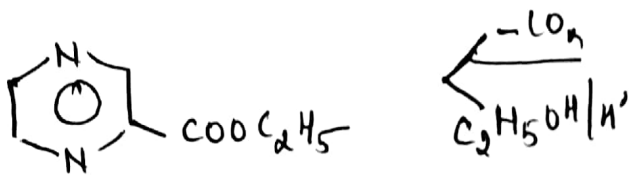
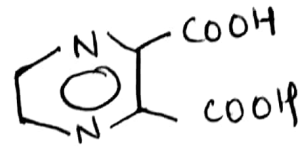
Pyrazine - 2. carboxamide

Syn :

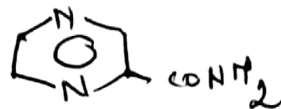


QUINOXALINE

$\downarrow$ (b)



$\downarrow \text{NH}_3$



(Pyrazinamide)

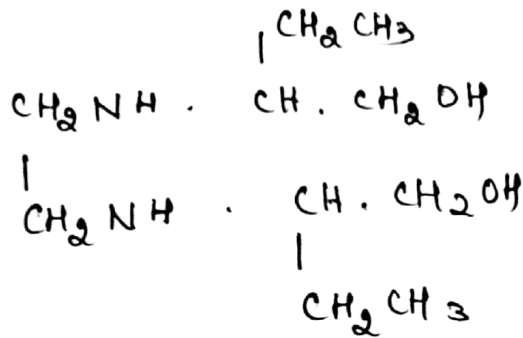
M.O.A :

Conversion of Pyrazinamide to pyrazinoic acid

Pyrazine carboxylic acid has bactericidal activity

To treat tuberculosis and Meningitis

ETHAMBUTOL



22' 1,2 Ethane diyl diamino bis-1-butanol HCl

M.O.A : Bacteriostatic drugs
inhibits the incorporation of mycolic acid
into the bacterial cell wall.

RIFAMPICIN

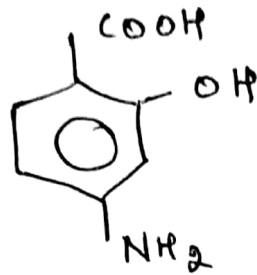
ANTIBIOTIC obtained from *Streptomyces*

mediterranei.

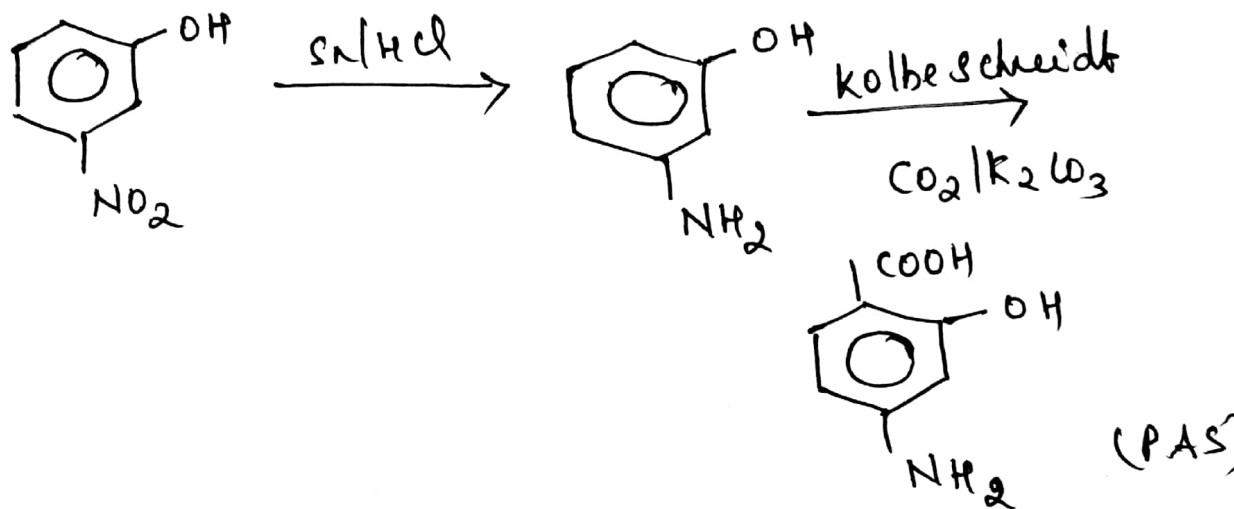
M.O.A : inhibits DNA dependant RNA
polymerase of mycobacteria
and inhibits the RNA synthesis and
act as a bactericidal drug

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PAS (Para Amino Salicylic acid)

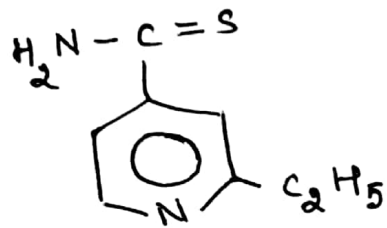


From m-nitrophenol

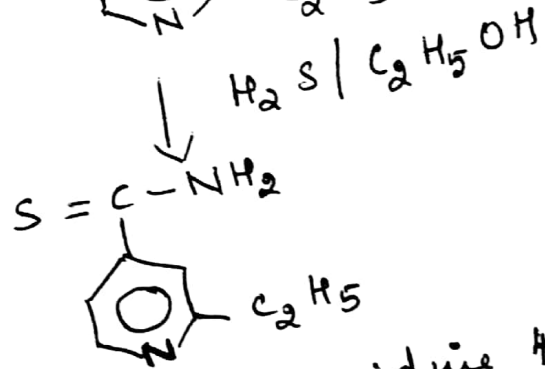
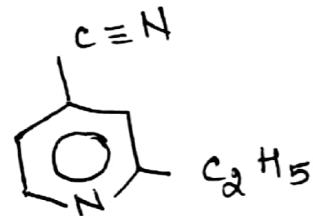
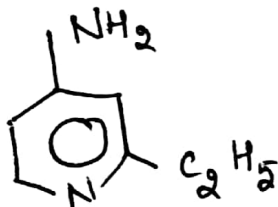


M.O.A . Antimetabolite

ETHIONANIDE



Synthesis



(2-Ethyl pyridine 4 carbothianide)

M.O.A :

Pro drug activated by an enzyme
Catalase peroxidase