

Indications: Has effect against a **variety of DNA viruses** – adenovirus, herpesviruses, papillomaviruses, polyomaviruses, & poxviruses

Most commonly used in treatment of **CMV retinitis**

Due to **significant nephrotoxicity**, usually used as a last resort agent in **drug resistant infections**

Mechanism of action: cidofovir is a **monophosphate nucleotide analogue**.

- Cellular uptake then cellular phosphorylation to active form **cidofovir diphosphate**. This has two actions:

- directly inhibiting viral DNA polymerase
- acts as alternate substrate for incorporation into newly formed DNA

These actions combine to result in growing **DNA chain termination**

Dose: IV only. Dosing frequency = weekly or fortnightly.

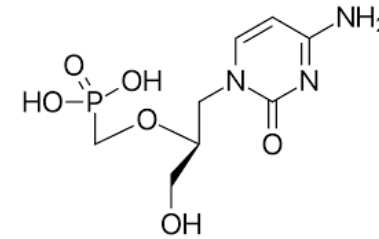
Must be given **with IV fluids before and after; AND probenecid**.

Probenecid provides **nephro-protection**. Cidofovir is primarily cleared by the kidneys. **Probenecid competes with Cidofovir** for tubular secretion. This results in maintaining the blood drug level in therapeutic range for longer, whilst reducing renal accumulation and nephrotoxicity.

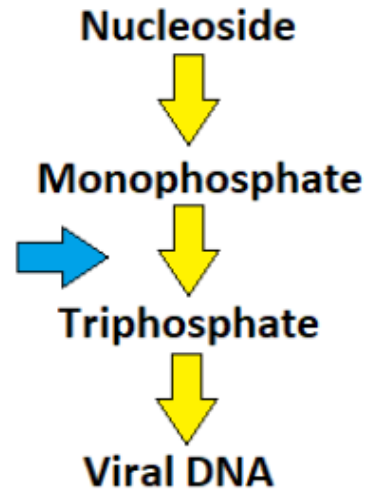
Adverse effects (please see BNF)

- nephrotoxicity – significant risk**. Usually reversible on discontinuation.
- neutropenia**

Cidofovir



Blocked by
Cidofovir



Use in Pregnancy/ Breast feeding
Lack of information available.
Manufacturer advises to avoid

Use in Renal impairment
Avoid in significant renal impairment