Mycophenolate Intravenous for Adults



Who can administer

May be administered by registered competent doctor or nurse/midwife

Important information

- Treatment with mycophenolate should be initiated and maintained by appropriately qualified specialists
- Imunosuppressants should be **continued in the peri-operative period** if being used for transplant patients ^(ref 1) For other indications, specialist review is indicated
- Supplies of this drug for IV use are routinely kept in stock in GUH
- See Monitoring requirements overleaf
- Flush before and after with glucose 5%
- Consider intravenous to oral switch as soon as possible as excellent bioavailability (see under dose for details)
- Some **clinically significant** drug interactions may occur, particularly with the use of other immunosuppressive therapy (check the manufacturers information carefully).
- IV to PO switch- see under Dose
- Brands are NOT interchangeable (see "Further information")

Available preparations

Cellcept 500mg vial

Reconstitution

Glucose 5%

14ml per 500mg vial - produces a 500mg in 15ml solution

Dilute further prior to administration

Infusion fluids

Glucose 5%

Methods of intravenous administration

Intermittent intravenous infusion (administer using an electronically controlled infusion device)

• Reconstitute as above- then dilute as follows to give a final concentration of ~ 6mg/ml

Dose	Add to this volume of infusion fluid
500mg (1 x 15ml)	70ml
1000mg (2 x 15ml)	140ml
1500mg (3 x 15ml)	210ml

Example: to prepare a 500mg dose, reconstitute a 500mg vial with 14ml Glucose 5%. Remove and discard 30ml from a 100ml Glucose 5% infusion bag (to make a 70ml volume), and add in the 15ml of reconstituted drug solution

Administer required dose over 2 hours, via either peripheral or central line

The residual volume in the infusion line must be flushed through at the same rate to avoid significant underdosing

Dose in adults

Usual dose

• Renal or hepatic transplantation (IV dose): 1g twice daily, started within 24 hours of transplantation and continued for 4 days in liver transplant, and up to a maximum of 14 days in either indication

Renal impairment

• Avoid doses greater than 1g twice daily in renal transplant patients with GFR < 25ml/min/1.73m² outside the immediate post-transplant period. Monitor the patient carefully.

Dose for Intravenous to Oral switch (ref 2)

- Cellcept = mycophenolate mofetil
- Myfortic = mycophenolic acid
- When switching from mycophenolate mofetil IV to oral mycophenolate mofetil, or vice versa, a dose adjustment is not needed (the manufacturers note a difference in doses of IV vs PO for hepatic transplant patients- but this is in the first 4 days post transplantation only)
- If switching from oral mycophenolic acid to IV mycophenolate mofetil- first convert the dose of mycophenolic acid to mycophenolate mofetil, and then change to IV

Monitoring

- FBC every week for 4 weeks, then twice a month for 2 months, then monthly in the first year.
- If **neutropenia** develops it may be appropriate to interrupt or discontinue treatment.
- Advise patient to report immediately any bleeding, bruising or infection which may indicate **bone marrow suppression**.

Further information

- **Avoid direct contact** of prepared solution with skin or mucous membranes: if such contact occurs wash thoroughly with soap and water. Rinse eyes with water.
- Avoid the use of live attenuated vaccines.
- Mycophenolate mofetil (Cellcept) is not interchangeable with mycophenolic acid (Myfortic). The
 brand of mycophenolate must be specified by the prescriber because of differences in bioavailability
 between brands.

Storage

Store below 25°C

References

SPC 16/02/2022

1: UKCPA The handbook for perioperative medicines, Transplant antirejection Medications

2:Uptodate downloaded 24/05/2022

Therapeutic classification

Immunosuppressant (purine synthesis inhibitor)