

Who can administer

May be administered by registered competent doctor or nurse/midwife

Important information

- Must be initiated and supervised by consultant neurologist
- In order to maintain traceability, the name and batch number of the administered product should be clearly recorded

Available preparations

Vyepti 100 mg per 1ml vial

Vyepti 300mg per 3ml vial

Reconstitution

Already in solution

Dilute further prior to administration

Infusion fluids

Sodium chloride 0.9%

Methods of intravenous administration

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

- Add required dose to 100mL Sodium chloride 0.9%
- Administer over 30 minutes

Dose in adults

Prophylaxis of migraine in adults who have at least 4 migraine days per month

- Recommended dose is 100mg every 12 weeks
- Some patients may benefit from a dose of 300mg every 12 weeks
- The need for dose escalation should be assessed within 12 weeks after initiation

Monitoring

- The patient should be monitored during and after the infusion in accordance with normal clinical practice
- Serious hypersensitivity reactions, including anaphylactic reactions, have been reported and may develop within minutes of the infusion.Â
- Please also consult Medication Protocol: Management of Infusion Related Patient Reactions in nurse-led infusion settings in GUHÂ [CLN-NM-0118](#)

Further information

- Overall benefit and continuation of treatment should be assessed 6 months after initiation of the treatment
- Contains sorbitol (E420). Patients with hereditary fructose intolerance (HFI) must not be given this medicinal product unless strictly necessary.

Storage

Store between 2 and 8°C

Do not freeze

References

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Therapeutic classification

CGRP receptor blocker