

Dosage



Recommended treatment:

3 months followed by 1 week drug-free to assess results. Repeat as long as symptoms persist.¹¹

INCREASING ONLY IF NEEDED

Tablets not actual size



1 x 120mcg
at bedtime

2 x 120mcg
at bedtime

1 x 240mcg

www.stopbedwetting.org

Prescribing Information: DesmoMelt® 120 and 240 micrograms oral lyophilisate.

Please consult the full Summaries of Product Characteristics before prescribing.

Name of Product: DesmoMelt 120 micrograms oral lyophilisate; DesmoMelt 240 micrograms oral lyophilisate. **Composition:** 120 or 240 micrograms of desmopressin (lyophilisate as acetate). **Indications:** Treatment of primary nocturnal enuresis (5 to 65 years of age). **Dosage and administration:** Children and adults (5–65 years of age) with normal urine concentrating ability: Initial dose of 120 micrograms sublingually at bedtime and if this dose is not sufficiently effective, the dose may be increased up to 240 micrograms, administered sublingually. Fluid restriction should be observed. DesmoMelt is intended for treatment periods of up to 3 months. The need for continued treatment should be reassessed by means of a period of at least 1 week without desmopressin. If adequate clinical effect is not achieved within 4 weeks following dose titration the medication should be discontinued. In the event of signs or symptoms of water retention and/or hyponatraemia treatment should be interrupted until the patient has fully recovered. When re-starting treatment fluid restriction should be enforced. **Contraindications:** Hypersensitivity to the active substance or to any of the excipients. Known or suspected cardiac insufficiency and conditions requiring treatment with diuretics, moderate and severe renal insufficiency. DesmoMelt should only be used in patients with normal blood pressure and they should not be used in patients with known hyponatraemia, syndrome of inappropriate ADH secretion or patients (SIADH) over the age of 65. Exclude diagnosis of psychogenic polydipsia (resulting in urine production exceeding 40 ml/kg/24 hours). **Side Effects:** Common: headache. Please consult the full Summary of product characteristics for further information about side effects. **Special Warnings:** Take care in patients with reduced renal function and/or cardiovascular disease or cystic fibrosis. In chronic renal disease the antidiuretic effect of DesmoMelt would be less than normal. Fluid intake must be limited to a minimum from 1 hour before until the next morning (at least) 8

hours after administration. Treatment without concomitant reduction of fluid intake may lead to water retention and/or hyponatraemia with or without accompanying signs and symptoms. All patients and, when applicable, their guardians should be carefully instructed to adhere to the fluid restrictions. **Precautions:** Severe bladder dysfunction and outlet obstruction should be considered before starting treatment. Elderly patients and patients with serum sodium levels in the lower range of normal may have an increased risk of hyponatraemia. Treatment with desmopressin should be interrupted during acute intercurrent illnesses characterised by fluid and/or electrolyte imbalance (such as systemic infections, fever, gastroenteritis). Caution should be used in: illnesses characterised by fluid and/or electrolyte imbalance; patients at risk for increased intracranial pressure. Hyponatraemia should be avoided by careful attention to fluid restriction and frequent sodium monitoring in case of concomitant treatment with drugs known to induce SIADH, treatment with NSAIDs and some antidiabetics of the sulfonylurea group particularly chlorpropamide. **Special precautions for storage:** None. **Presentation:** Carton containing 30 oral lyophilisates in blister strips. **Marketing Authorisation Number:** 120 micrograms 03194/0094, 240 micrograms 03194/0095. **Marketing Authorisation Holder:** Ferring Pharmaceuticals Ltd., Drayton Hall, Church Road, West Drayton, UB7 7PS. **Legal Category:** POM. **Basic NHS Prices:** 30 x 120 micrograms £30.34, 30 x 240 micrograms £60.68. **Date of Preparation:** December 2019. All trademarks registered to Ferring. **PI approval code:** UK-MN-1900002

Adverse events should be reported. Reporting forms and information can be found at www.mhra.gov.uk/yellowcard. Adverse events should also be reported to Ferring Pharmaceuticals Ltd. Tel: 0844 9310050. Email: medical.uk@ferring.com

References

- Hjälmsås K. *Acta Paediatr* 1997;**86**(9):919–22.
- Yeung CK et al. *BJU Int* 2006;**97**(5):1069–73.
- Janknegt RA et al. *J Urol* 1997;**157**(2):513–7.
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- van Kerrebroeck PEV. *BJU Int* 2002;**89**(4):420–5.
- Warzak WJ. *Clinical Pediatrics* 1993;**32**:38–40.
- Lottman H et al. *Int J Clin Pract* 2007;**61**(9):1454–60.
- Debruyne P et al. Poster presented at ICCS/ERIC/BAJU Joint Congress in London, UK, 12–14 October 2012.
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- NICE costing statement 2010. Nocturnal Enuresis.
- DesmoMelt Summary of Product Characteristics (SmPC).

Understanding the true value of being
home and dry



DESMOMELT®
Desmopressin (as acetate)
120mcg and 240mcg oral lyophilisate



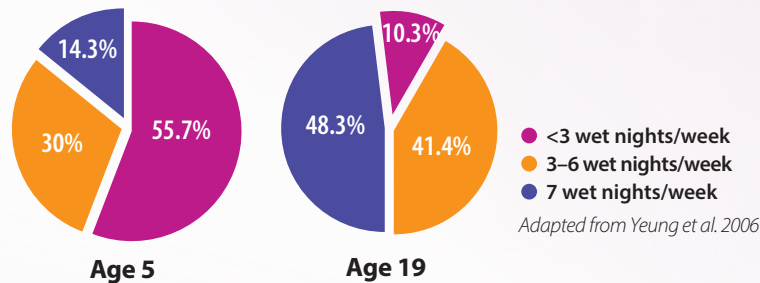
Primary Nocturnal Enuresis (PNE)

Primary Nocturnal Enuresis (PNE) is widespread, and is the most common chronic ailment in children besides allergic disorders¹ affecting approximately 16% of 5 year old children²

PNE is a distressing condition that can have a significant impact on a young person's behaviour and their emotional and social wellbeing³

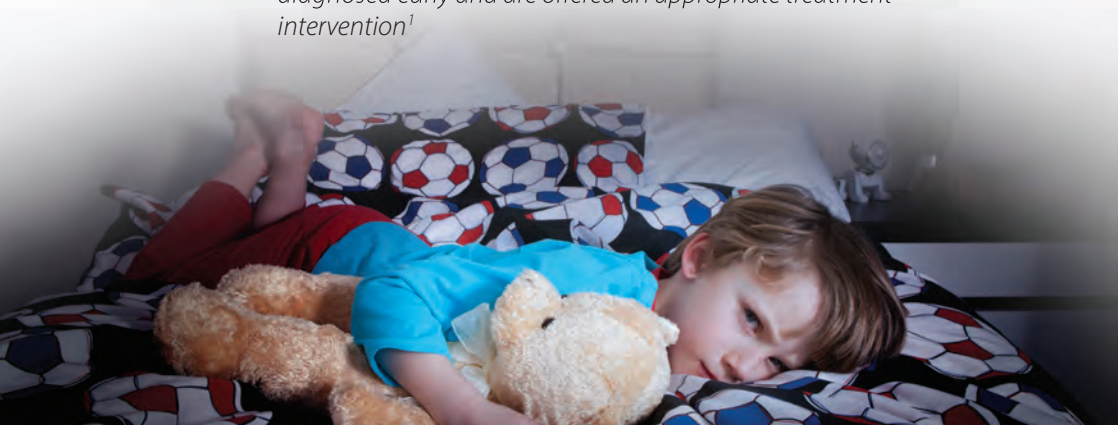
- 32.5% of children (aged 9) rated wetting the bed as a 'quite difficult' life event for them⁴
- They rated bedwetting to be comparable to how they felt about being teased or not being able to spell properly⁴

PNE can be a persistent problem, and 10% of enuretic children will remain bedwetters for life¹



The burden of PNE on children and their families is considerable,⁵ however, effective treatment benefits both parties⁶

- So, it is important that children suffering with PNE are diagnosed early and are offered an appropriate treatment intervention¹



Efficacy



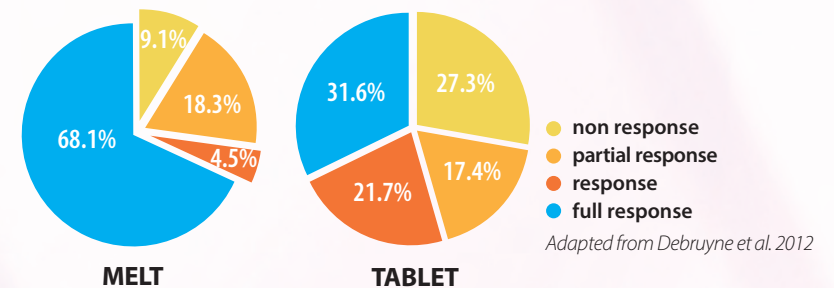
DesmoMelt is well accepted by children of various ages and facilitates early intervention⁷

DesmoMelt provides an effective and convenient solution to PNE, helping children and families return to a normal life

- Statistically significant preference for **DesmoMelt** compared with tablets in children aged 5–11 years⁷

DesmoMelt is associated with higher compliance than tablets and retains similar levels of efficacy and safety at lower dosing levels⁷

- High compliance of **94.5%**⁷
- In a study comparing children's clinical response rates after 2 months treatment, overall response rates were better with **DesmoMelt**⁸



Understanding the true value of DesmoMelt

- **Statistically significant** preference for **DesmoMelt** compared with tablets in children aged 5–11 years⁷
- **Statistically significant superior** antidiuretic effect at 3–8 hours after dosing when compared to tablet⁹
- **The value of compliance**⁷ with no significant cost burden to NHS¹⁰