



## Final Appraisal Recommendation

**Alteplase (Actilyse<sup>®</sup> Cathflo<sup>®</sup> 2 mg)**

**Limited submission by:  
Boehringer Ingelheim Ltd**

**Advice No: 1911 – December 2011**

### Recommendation of AWMSG

**Alteplase (Actilyse<sup>®</sup> Cathflo<sup>®</sup> 2 mg) is recommended as an option for use within NHS Wales for the thrombolytic treatment of occluded central venous access devices including those used for haemodialysis.**

**AWMSG is of the opinion that alteplase (Actilyse<sup>®</sup> Cathflo<sup>®</sup> 2 mg) is not suitable for shared care within NHS Wales.**

#### **Additional notes:**

This recommendation applies only to the use of alteplase 2 mg vial (Actilyse<sup>®</sup> Cathflo<sup>®</sup> 2 mg) for the indication under consideration. The alteplase 2 mg vial (Actilyse<sup>®</sup> Cathflo<sup>®</sup> 2 mg) is the only presentation of alteplase licensed for use in this indication.

Alteplase (Actilyse<sup>®</sup> Cathflo<sup>®</sup> 2 mg) for the above indication met the following criteria for eligibility for a limited submission:

- New formulation with a pro-rata or lower cost per treatment.
- Anticipated usage in NHS Wales is considered to be of minimal budgetary impact.

In reaching this recommendation AWMSG took account of the [AWMSG Secretariat Assessment Report](#), the preliminary appraisal recommendation (PAR) and the applicant company's response to the PAR, lay perspective and discussions at AWMSG.

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