



Final Appraisal Recommendation

Advice Number: 0319 – March 2019

**Dasatinib (Sprycel®) 20 mg, 50 mg, 70 mg, 80 mg, 100 mg and
140 mg film-coated tablets**

**Limited submission by Bristol-Myers Squibb Pharmaceuticals
Ltd**

**Recommendation of the All Wales Medicines Strategy
Group**

Dasatinib (Sprycel®) is recommended as an option for use within NHS Wales for the treatment of paediatric patients weighing ≥ 10 kg with newly diagnosed Philadelphia chromosome positive (Ph+) chronic myelogenous leukaemia (CML) in chronic phase (Ph+CML-CP) or Ph+CML-CP resistant or intolerant to prior therapy including imatinib.

This recommendation applies only in circumstances where the approved Patient Access Scheme (PAS) is utilised or where the list/contract price is equivalent to or lower than the PAS price.

Additional note(s):

- Please refer to the Summary of Product Characteristics for the full licensed indication.
- AWMSG considered that dasatinib (Sprycel®) satisfied the AWMSG criteria for a medicine developed specifically for rare diseases, as an orphan equivalent.
- The 70 mg oral tablet is not currently marketed in the UK.

In reaching the above recommendation AWMSG has taken account of the appraisal documentation prepared by the AWMSG Secretariat (reference number 1514), which includes the AWMSG Secretariat Assessment Report (ASAR), the Preliminary Appraisal Recommendation (PAR) and the applicant company's response to the PAR, clinical expert opinion (where available), the views of patients/patient carers (where available) and the lay member perspective.

The All Wales Therapeutics and Toxicology Centre (AWTTC) reviewed this appraisal recommendation in June 2022. It will be reviewed again in three years.

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