Ivacaftor for the treatment of patients with cystic fibrosis and the G551D mutation: a systematic review and cost-effectiveness analysis


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Authors' objectives
To review the clinical effectiveness and cost-effectiveness of ivacaftor for the treatment of CF in patients aged 6 years who have the G551D mutation.

Authors' conclusions
The available evidence suggests that ivacaftor is a clinically effective treatment for patients with CF and the G551D mutation; the high cost of ivacaftor may prove an obstacle in the uptake of this treatment. The main priority for further research is the long-term effectiveness of ivacaftor.

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