cullect				
Study	[Chalayer 2016 ¹⁶⁵]			
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness
Economic analysis: CUA (health outcome: QALYs) Study design: Decision analytic model Approach to analysis: A decision tree based on results of Palumbo 2011 clinical trial ⁷²⁴ .	Population: Patients newly diagnosed with multiple myeloma treated with protocols including thalidomide Cohort settings: Start age: NR Male: NR	Total costs (mean per patient): Intervention 1: £230 Intervention 2: £1,283 Incremental (2–1): £1,053 (95% CI: NR; p=NR) Currency & cost year: 2013 Euros (presented here as 2013 UK pounds ^(b))	QALYs (mean per patient): Intervention 1: 0.300 Intervention 2: 0.299 Incremental (2–1): -0.001 (95% CI: NR; p=NR)	ICER (Intervention 2 versus Intervention 1): Intervention 1 dominant (less costly and more effective)(pa) 95% Cl: n/a Probability Intervention 2 cost-effective (£20K/30K threshold): NR Analysis of uncertainty: None of the sensitivity analyses undertaken changed the conclusion.
Perspective: France National Health Insurance System Time horizon: 6 months Treatment effect duration: ^(a) 6 months Discounting: Costs: n/a ; Outcomes: n/a	Intervention 1: Aspirin (100mg/day) for 3 months. Intervention 2: LMWH standard dose, standard duration) (Enoxaparin 40mg/day) for 6 months.	Cost components incorporated: Hospitalisation GP visits Home nursing Laboratory investigation Radiologic procedures Drugs		

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Cancer

Health outcomes: data on baseline risks and relative treatment effects are based on a single RCT (Palumbo 2011⁷²⁴). These outcomes included DVT, PE, stroke, acute MI, major bleeding and sudden death. Quality-of-life weights: EQ-5D index values were used. Cost sources: National unit cost sources were used including National reimbursement database and Vidal drug compendium.

Comments

Source of funding: None. Limitations: Some uncertainty regarding the applicability of unit costs from France in 2013 to current NHS context. The model does not incorporate any long-term consequences such as CTEPH or PTS. Baseline risk and relative treatment effects are based on a single open-label trial, so by definition, does not reflect all available evidence. Costs of LMWH administration might be underestimated.

Overall applicability:^(c) Partially applicable **Overall quality**^(d) potentially serious limitations

Abbreviations: 95% CI: 95% confidence interval; CTEPH: chronic thromboembolic pulmonary hypertension; CUA: cost-utility analysis; da: deterministic analysis; EQ-5D: Euroqol 5 dimensions (scale: 0.0 [death] to 1.0 [full health], negative values mean worse than death); ICER: incremental cost-effectiveness ratio; n/a: not applicable; NR: not reported; pa: probabilistic analysis; PTS: post-thrombotic syndrome; QALYs: quality-adjusted life years

- (a) For studies where the time horizon is longer than the treatment duration, an assumption needs to be made about the continuation of the study effect. For example, does a difference in utility between groups during treatment continue beyond the end of treatment and if so for how long.
- (b) Converted using 2013 purchasing power parities⁷¹⁵
- (c) Directly applicable / Partially applicable / Not applicable
- (d) Minor limitations / Potentially serious limitations / Very serious limitations