A238 Abstracts

ES₅

Stroke-Eprosartan Compared to Nitrendipine for Secondary Prevention'). Costs and utilities were derived from published estimates considering country-specific health care payer perspectives. Drug prices of the comparators were based on the cheapest generics. The treatment time horizon simulated was 2.5 years (mean observation period of the MOSES study) modelling follow-up effects over a life-time. Costs and effects were discounted according to country-specific guidelines. RESULTS: During a 2.5-year treatment time horizon modelling follow-up effects over lifetime in 1,000 patients eprosartan treatment avoided about 58 events (26 cardiovascular, 32 cerebrovascular) and produced about 30 incremental QALYs versus the compared treatments. Irrespective of country and comparator the cost per QALY gained by eprosartan never exceeded €20,000 and therefore went far below the estimated willingness-to-pay threshold of €30,000. The probabilistic sensitivity analyses fortify these outcomes by showing a probability of 90-100% that eprosartan is a cost-effective treatment strategy. CONCLUSION: Even comparing eprosartan to low-priced generic substances, the HEALTH model simulations provide evidence that eprosartan treatment is associated with obvious health benefits being obtained at reasonable cost. Eprosartan should therefore be considered a good treatment option for hypertensive patients with cerebrovascular disease.

PODIUM SESSION IV: ECONOMIC STUDIES II

DOES ARTHROSCOPIC ACROMIOPLASTY PROVIDE ANY ADDITIONAL VALUE IN THE TREATMENT OF SHOULDER IMPINGEMENT SYNDROME? A TWO-YEAR RANDOMIZED CONTROLLED TRIAL

Ketola S¹, Lehtinen J², Arnala I¹, Nissinen M¹, Westenius H¹, Aronen P³, <u>Sintonen H³</u>, Konttinen Y³, <u>Malmivaara A⁴</u>, Rousi T¹ ¹Kanta-Häme Central Hospital, Hameenlinna, Finland, ²Tampere University, Tampere, Finland, ³University of Helsinki, Helsinki, Finland, ⁴Finnish Office for Health Care Technology Assessment, Helsinki, Finland

OBJECTIVES: To examine in randomized controlled trial the effectiveness and cost-effectiveness of arthroscopic acromioplasty in the treatment of stage II shoulder impingement syndrome. METHODS: We divided 140 patients into supervised exercise program (n = 70, exercise group) and arthroscopic acromioplasty, followed by a similar exercise program (n = 70, combined treatment group). The primary health outcome measure was self-reported pain on a 0-10 Visual Analogue Scale at 24 months with a two-point change defined as minimal clinically important difference (MCID). RESULTS: Results In an intention-to-treat analysis an improvement exceeding MCID took place from baseline to 24 months in both groups: self-reported pain diminished from 6.5 to 2.9 in the exercise group (N = 66) and from 6.4 to 2.5 in the combined treatment group (n = 68) (P < 0.001 in both). In the combined treatment group pain relief was attained faster, but the groups did not any more differ at 24 months (P = 0.37). A similar pattern was seen in the secondary outcome measures: disability, pain at night, SDQ score, ability to work, number of painful days and proportion of pain-free patients. The mean total cost was €2961 in the combined treatment group and €1864 in the exercise group. The incremental cost-effectiveness ratio was €5852 per MCID unit, i.e., combined treatment was considerably more costly. CONCLUSION: Arthroscopic acromioplasty does not provide any significant additional value over structured and supervised exercise program alone in terms of subjective outcome or cost-effectiveness. Operative treatment should be offered judiciously.

ES6

COST COMPARISON BETWEEN HAEMODIALYSIS AND PERITONEAL DIALYSIS IN NORWAY FOR PATIENTS WHO CAN USE EITHER TREATMENT MODALITY

Nyhus K, Kristensen FKO, Merméjean P, Sverre JM PharmEcon, Asker, Akershus, Norway

OBJECTIVES: For patients with renal failure who from a medical perspective can be treated with either haemodialysis (HD) or peritoneal dialysis (PD), the choice of treatment is mainly based on administrative, economic and patient preference considerations. HD is performed 3-5 times per week in a hospital, while PD is performed daily at home. Both are financed over hospitals' budgets in Norway. This evaluation compared the costs of HD and PD from both a societal and a hospital perspective for such patients. METHODS: Costs were calculated based on national data on resource use and unit costs. Estimates of resource use were based on treatment guidelines, the literature and interviews at three major hospitals in Norway. Unit costs were based on tariffs, price lists, hospital accounts and salary statistics. In the societal perspective, costs were divided in three sections: Costs born by patients (co-payments, transportation and value of time), costs born by hospitals (personnel, medicines and supplies, laboratory tests, capital and infrastructure) and other public costs (funding of medicines). In the hospital perspective, net cost was calculated (difference between expenses and income). RESULTS: From a societal perspective the average monthly cost per patient is for PD: 30,700 NOK (~3,800€) and for HD: 51,800 NOK (~6,400€). The main cost driver for PD is dialysis solutions: 23,800 NOK (~2,900€). The cost components in HD are more homogeneous. Hospitals go about break-even performing PD and have a monthly net income of about 600 NOK (~75€) when performing HD. CONCLUSION: For patients who can use either treatment modality, PD is a cost saving alternative to HD from a societal perspective. However, the financing systems for dialysis in Norway make hospitals relatively neutral from an economical perspective in their choice of HD or PD for these patients.

ES7

ASSESSMENT OF LONG-RUN ECONOMIC BENEFITS ASSOCIATED WITH IN-VITRO FERTILIZATION (IVF) FUNDING DECISIONS: A SIMPLIFIED LIFETIME TAX CALCULATION

 $\underline{\text{Connolly MP}}^{I}, \text{Hoorens S}^{2}, \text{Gallo F}^{2}, \text{Ledger WL}^{3}$

¹Ferring International Center, Saint-Prex, Switzerland, ²RAND Europe, Cambridge, UK, ³University of Sheffield, Sheffield, UK

OBJECTIVES: Globally there is considerable variation in public funding for IVF treatments. IVF is unique amongst health interventions because its success leads to human life. In light of this uniqueness we apply a Generational Accounting approach, an accepted method used by tax authorities, to assess whether publicly funded IVF represents sound fiscal policy. Our assessment considers future lifetime net tax contributions to the British government (taxes paid minus transfer payments) attributed to a successful IVF birth. METHODS: Net present value (NPV) calculations were applied to the average cost per successful IVF conceived live birth (£12,931 in 2005), lifetime direct cash transfers and lifetime future tax contributions discounted using established Treasury department rates. We assume the following: full-time education aged 6-19; full-time employment aged 20-68 (Pension Commission, 2005); education costs, child-tax credits, and pension contributions increase with inflation. Age-specific income was adjusted for inflation over time, and we allowed expected income to vary with age. Current government tax revenues of 35.5% gross income were held constant. RESULTS: Based on average life-expectancy the model indicates an