NOT ALL PATIENTS ARE AVERAGE: THE IMPORTANCE OF RECOGNISING PATIENT HETEROGENEITY
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OBJECTIVES: Cost-effectiveness analyses are routinely based on data from group averages, restricting its generalizability to those with below- or above-average risk. A pharmaeco-economic model was developed that used individualised risks, taking as example bisphosphonates and prevention of fractures. METHODS: Data were obtained from a research database of general practitioners, comprising a sample of the UK general population of women >50 years (N = 330,000). Individual mortality and hip, vertebral, and other osteoporotic fracture risks were estimated by age, sex, body mass index, smoking and other clinical risk factors. Estimates on costs, EQ5D utilities and treatment efficacy were obtained from a UK national report (NICE) and outcomes were simulated over a ten-year period. RESULTS: There was a large variability in the cost-effectiveness with clinical risk factors. At age 60–69, the cost per QALY gained was $136k in women with low fracture risk but $36k with high fracture risk (data for women without fracture history). Patients with low body mass index (<20) had considerable better cost-effectiveness than patients with high BMI (≥26) ($23k versus $71k at age 60–79 in women without fracture history). The same was found for different diseases such as rheumatoid arthritis or inflammatory bowel disease. Using a cost-acceptability ratio of $30k per QALY gained, bisphosphonates became cost-effective for patients with a 5-year risk of 9.3% (95% CI 8.0–10.5%) for osteoporotic fractures and of 2.1% (95% CI 1.5–2.7%) for hip fractures. Including bone mineral density in the risk assessment, the cost per QALY gained was $35k in women at age 60 with a fracture history and a T-score of -2.5 (at age 80, this was ≤3k). CONCLUSIONS: A pharmaeco-economic model based on individual long-term risks (as derived from a health care database) can improve the targeting in a cost-effective manner of therapy to patients.

BAYESIAN ESTIMATION OF COST-EFFECTIVENESS ADJUSTED FOR REAL WORLD CONSIDERING STATISTICAL ERRORS IN CLINICAL TRIALS
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Although evidence-based treatment is a norm for practice, the efficacy of treatment presented in clinical trials does not necessarily guarantee its full size of benefit in a real world. One of the factors to be considered is statistical error, which may cause a discrepancy between “efficacy” in a clinical trial and “effectiveness” in real practice. OBJECTIVES: The purpose of this study is to formulate the influence on the cost-effectiveness in practice caused by statistical errors when a treatment is undertaken on the basis of the evidence proven in a clinical trial. METHODS: A decision analysis was performed by decision-tree modeling with treatment options that reflected the alternatives: take a new treatment A (TA) vs. a conventional treatment B (TB), depending on the evidence resulted from a clinical trial. The rule for selecting a treatment is: 1) take TA if the clinical trial confirmed better efficacy with TA than TB, or 2) keep using TB if failed to prove dominancy of TA over TB. Also the decision tree modeled Bayesian prior probabilities at the chance node in the first depth with both the null hypothesis H0 and the alternative H1 (in efficacy, TA = TB, and TA > TB, respectively). RESULTS: Folding-back and averaging-out procedures in the decision-tree led to a mathematical formula with the parameters such as cost-effectiveness ratios, type I and type II errors, and Bayesian prior probabilities for treatment effect. Numerical analysis for the developed formula delineated an expected decrease of cost-effectiveness is not always negligible depending on the statistical power of the clinical trial and also on the degree of prior belief in a new treatment from the standpoint of Bayes. CONCLUSIONS: In case of need for adjusting cost-effectiveness from clinical trial into real practice, the formula newly developed can be a tool to measure the degree of the adjustment.

MEASURING SOCIAL PREFERENCES FOR EQ-5D HEALTH STATES: NEW SOLUTIONS TO OLD PROBLEMS
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OBJECTIVES: The standard version of EQ-5D defines 243 health states. Eliciting corresponding social preference weights is a labour/resource intensive exercise—witness the scale of the original 1993 UK valuation study and the subsequent US replication study. This stems from the requirement to use TTO methods in face-face interviews. Furthermore, given the need to minimise respondent burden only a subset of 45 health states are directly valued. These data are used to construct an estimation model used to interpolate values for the remaining health states. The complexity of this modelling represents a daunting challenge. Simpler methods are called for. This paper reports on a study in which values for all EQ-5D health states were obtained directly. METHODS: The EuroQoL Group has a standard questionnaire for use in collecting valuations in postal surveys. Multiple versions of this questionnaire were designed, each presenting a subset of 16 EQ-5D health states drawn from across the severity range. Values for health states are assessed on a 0–100 scale that represents worst-best imaginable health. A total of 1100 questionnaires were mailed to a sample of respondents selected from the electoral registers of England and Wales. Data from 685 respondents were subsequently analysed. RESULTS: OLS regression was used to determine the value decrements associated with increased problems on each EQ-5D dimension. The adjusted R² for this model is 0.907. Mean values from this survey were compared with mean VAS scores elicited in the original 1993 survey. For 30/45 health states values were remarkably similar and differed by less than 5 points. The highest proportional difference occurs for dead with a value in 2003 that is 45% higher than that recorded in 1993. CONCLUSIONS: The direct valuation of EQ-5D health states is feasible and yields robust results that avoid the use of problematic TTO procedures.

CONVERTING SF-36 INTO EQ-5D: DESIGNED TO SUCCEED OR CALCULATED TO FAIL?
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BACKGROUND: In its original format SF-36 is a profile measure that results in 2 or more subscale scores. This design means that SF-36 cannot be used in cost-effectiveness analysis in which health outcomes must be represented as a single summary index of the type exemplified by EQ-5D utility. However, many clinical studies continue to report outcomes in terms of SF-36 and the researchers continue to confront the challenge of “salvaging” such data for use in economic analysis. This paper reviews the dangers in using SF-36/EQ-5D conversion models that have been
META-ANALYSIS OF ANTIDEPRESSANT TREATMENT EFFECTS IN PATIENTS AFTER STROKE

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OBJECTIVE: To systematically evaluate and assess the effectiveness of antidepressants in treating depressed patients after stroke. METHODS: A systematic review with Meta analysis of double-blinded randomized clinical trials (RCTs) about the treatment of depression in the patients with stroke. Data were collected from the published RCTs, which indexed in the Medline from 1960 to 2005. Review was only limited in the English articles. Outcome measurements: 1) The proportion of treatment responders whose depression scores improved by >50% was compared. 2) The depression rating scores at the beginning, middle, and end of treatments. RESULTS: A total of 313 patients from 5 RCTs fulfilling the inclusion criteria were identified for this study. The percentages of responders in treatment and control group were 57.9%, and 44.4%, respectively (Risk Difference, RD = 0.17, 95%CI from -0.04 to 0.38, Z = 1.61, P = 0.11). In term of depression scores, prior to any treatments, the overall depression scores in the patients of treatment groups were a little bit higher (more depressed) than the patients in the control groups, but the difference was not statistically significant (Weighted Mean Difference, WMD = 1.01, Z = 1.59, P = 0.11). After the treatments, patients in the treatment groups became statistically less depressed than the patients in the control groups in term of lower depression scores (WMD = -1.88, Z = 2.16, P = 0.03). In addition, we found that the overall WMD in term of depression scores between two groups was 0.60 (Z = 0.87, p = 0.39) at the beginning of treatment, and became -1.85 (Z = 2.12, P = 0.03) in the middle of treatment, then further decreased to -3.06 (Z = 2.92, p = 0.004) at the end of treatment. CONCLUSION: The results indicated that antidepressant treatments were effective in the patients after stroke in term of reducing the depression scores and treatment effects were time dependent. In addition, more efforts should be guaranteed to study the cerebrovascular effects of antidepressants in patients after stroke.

AUSTRALIAN SCHIZOPHRENIC PATIENTS TREATED WITH RISPERIDONE LONG-ACTING INJECTION (RLAI): INTERIM RESULTS FROM THE E-STAR STUDY

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OBJECTIVE: To determine time to discontinuation in schizophrenic patients who commence RLAI. METHODS: e-STAR (electronic-Schizophrenia Treatment Adherence Registry) is an ongoing international observational study of schizophrenic patients who commence RLAI. Data collected retrospectively (12-months) and prospectively (2-years) included: patient demographics, medications, hospitalisations, Clinical Global Impression-Severity (CGI-S), Global Assessment of Functioning (GAF) and adverse events. A priori statistical analyses using Kaplan-Meier curves and proportional hazard regression models included time to discontinuation. RESULTS: Data from 591 patients from 15 hospitals were available for this analysis. The average age was 36.9 [12.5] years (mean [SD]), duration of illness was 11.0 [9.3] years, 71% were male. The baseline CGI-S was 4.5 [1.2] and GAF was 41.8 [14.8]. Ninety-one percent of