

Personal Social Services (PSS). Costs were calculated based on standard United Kingdom sources.

RESULTS:

The estimated incremental cost-effectiveness (ICER) of dexamethasone compared with LCP was GBP19,509 per quality-adjusted life year (QALY) gained. The estimated ICER of adalimumab compared with LCP was GBP94,523 and GBP317,547 per QALY in patients with active and inactive uveitis respectively. The factors with the largest impact upon the ICERs were the rate of blindness and the proportion of cases of blindness avoided by interventions.

CONCLUSIONS:

Dexamethasone and adalimumab resulted in health gains, but at significant extra costs, especially adalimumab which is unlikely to be considered a cost-effective use of NHS resources. The results of the analysis are highly uncertain due to the limited availability of evidence on: the comparative effectiveness of dexamethasone, adalimumab and current practice; the effectiveness of treatments in avoiding blindness; and, the effectiveness of interventions in different subgroups.

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OP132 How A Shared Management Of Home Infusion Can Control Expenditure

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INTRODUCTION:

In France, medical devices (MDs) for home-based infusion used to be covered by the health insurance system if included on the list of products and services qualifying for reimbursement under a generic description corresponding to a class of products with the same indications. This coverage modality offered no resistance to unnecessary or wasteful spending. Besides, between 2010 and 2015 the expenditure related to these MDs have increased from EUR192million to EUR289million (+50 percent).

METHODS:

The French National Authority for Health (HAS) has assessed the actual benefit of these MDs which have the same indications as the drugs infused at home. This work led to standardize the infusion types (gravity, elastomeric pump or active system requiring an energy source) and the quantities of MDs needed to carry out the different cares (installation, connection, withdrawal) according to the infusion route. At this step, considering that the priority was to redefine the MDs required at home for each care type, no economical assessment had been conducted.

RESULTS:

Based on this medical assessment, the Ministry of Health has distinguished three types of infusion and three types of services (home installation, monitoring and consumables) since 2016. In total, twenty-four packages have been set up for reimbursement with non-cumulative rules. Doctors are in charge to prescribe the appropriate packages; providers and nurses determine together the optimal devices needed for each patient according to his environment.

CONCLUSIONS:

These HAS recommendations on practice standardization have been the keystone for cost negotiations. The new coverage modalities aim to motivate liberal nurses to choose the best fitted products and providers to deliver the right quantities to patients. The expected benefits are an adjusted evaluation of the necessary equipment and a control of health expenditure due to the fixed costs of each infusion package.

OP133 Health Technology Assessment In Brazil: A 5-year Review Of Brazilian Health System (CONITEC) Activities

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INTRODUCTION:

Since the creation of the National Committee for Health Technology Incorporation in the Brazilian Health System (CONITEC), a new phase started in the public Brazilian Health System (SUS): a continuous updating of the system based on Health Technology Assessment (HTA). CONITEC was created by federal law in 2012 and is responsible for advising the Ministry of Health regarding the incorporation or disinvestment of health technologies. The whole process involves a strong interaction with society, including the composition of the committee, which has the participation of the National Health Council. The objective of this study was to describe the results of CONITEC in five years of operation.

METHODS:

This is a retrospective descriptive study, based on information from the database (period 2012–2016) and CONITEC's website.

RESULTS:

Since 2012, CONITEC assessed 541 technologies, including drugs (360), health products (71) and procedures (110); 303 assessment requests came from SUS agencies and institutions and the other 238 requests from pharmaceutical companies, medical societies, patient associations and the judiciary bodies. In this period, there were 190 public consultations, during which more than 24,000 feedback from society were received. The average time for evaluation was 146 days. The committee recommended the incorporation of 186 technologies into SUS, the disinvestment of 43 and was unfavorable to the incorporation of 88, generating a budgetary impact of approximately BRL2.5 billion (USD764 million).

CONCLUSIONS:

From 2012–2016, CONITEC tripled the average annual incorporation of new technologies compared to the period 2006–2011. In this process, it was necessary to assess efficacy, safety and cost-effectiveness of technologies, generating positive results for the expansion of access, health gains for patients and sustainability for the system. It should be considered that the use of evidence for decision making strengthens transparency in public management and the development of active processes of information, communication and social participation.

OP134 Predictors Of Public Health Outcomes: A Case Study From Turkey

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INTRODUCTION:

In Turkey, there is a scarcity of knowledge about the predictors of health outcomes at a national level, and it is well known that there is a gap between rural and urban parts of developing countries in terms of the level
