# OP127 Analysis Of The Competencies To Be Acquired In Health Technology Assessment

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## **INTRODUCTION:**

Health Technology Assessment (HTA) is a multidisciplinary activity that systematically examines different dimensions related to the direct and indirect consequences of health technologies when implemented in healthcare systems. HTA is developed by multidisciplinary teams in order to cover all the mentioned dimensions. However, the quality of the processes produced by HTA teams will depend upon the competencies that those teams will acquire and incorporate including knowledge, skills and attitudes (1). The aim of this research was to determine how well these dimensions and competencies are covered in HTA academic Masters degree courses and manuals.

## **METHODS:**

We analyzed what had been done in terms of competencies definition in HTA: how it has been reflected; theoretically and according to the authors, and how competencies can be structured; know-how and values. We explored HTA manuals and HTA academic Masters degree courses. We searched in Google with specific terms: building capacities, HTA, programs, Masters, diplomas. We used the HTAi vortal and the information related to courses (for example Masters degrees) and HTA agencies and network webpages for programs. The inclusion criteria were formal programs that describe HTA capacity building and not partial teaching of certain aspects of HTA and we excluded non-recognized institutions, or where there was no description of the programs or lack of detail regarding objectives and competencies to be achieved.

#### **RESULTS:**

We found 105 courses or programs and analyzed 8 reports and 3 manuals. The main challenges that we faced were: that information was difficult to retrieve, not similarly structured, difficulties to find key information in webpages, no program description at all in some cases and the need to contact institutions staff or register as a student to receive the information and finally, it was difficult to obtain contact details of key people. We structured the information on competencies in knowledge, skills and attitudes.

#### **CONCLUSIONS:**

The analyzed Masters degree courses and manuals did not cover all of the dimensions of HTA analysis in an equal and standardized way. The ethical, legal, social and organizational aspects were lacking in some of the programs, while, on the contrary, clinical and economic aspects were substantially included. On the basis of the information retrieved it would be good to define core competencies for HTA.

#### **REFERENCE:**

1. Mueller D, Gutiérrez-Ibarluzea I, Schuller T, et al. Capacity building in agencies for efficient and effective health technology assessment. *Int J Technol Assess Health Care*. 2016;32(4):292-299.

# OP129 Predictors Of Effectiveness In Patients With Rheumatoid Arthritis

#### **AUTHORS:**

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#### **INTRODUCTION:**

Biological disease-modifying anti-rheumatic drugs (bDMARDs) have become firmly established in the management of patients with rheumatoid arthritis (RA),

but some patients do not improve despite therapy. This study evaluated the predictors of effectiveness of the bDMARDs on a cohort of patients with rheumatoid arthritis (RA) in the Brazilian Public Health System.

#### **METHODS:**

RA individuals treated with bDMARDs, were included in the open prospective cohort study. The Clinical Disease Activity Index (CDAI) was used to assess the effectiveness comparing results at baseline and after 6 months of follow-up. The association between socio-demographic and clinical characteristics with the disease activity measured by the CDAI was also investigated. The bDMARDs was considered effective when the patient achieved remission or low disease activity and considered not effective when there was still moderate or high disease activity. Pearson's chi-square was applied for the univariate analysis to evaluate the association of effectiveness measured by the CDAI with the socio-demographic (gender, education, marital status and race) and clinical variables (type of drug, EuroQol (EQ)-5D and Health Assessment Questionnaire (HAQ)). Logistic regression was applied in the multivariate analysis of the variables that presented a p < .20 value during the univariate analysis.

### **RESULTS:**

All 266 RA patients completed six months of follow-up. The most widely used bDMARDs was adalimumab (57.1 percent), with etanercept used by 22.2 percent, golimumab by 7.5 percent, abatacept by 4.5 percent, tocilizumab by 3.4 percent, infliximab by 2.6 percent, certolizumab by 1.5 percent, and rituximab by 1.1 percent. The bDMARDs reduced disease activity as measured by CDAI at six months of follow-up (p<.001). The percentage of patients achieving remission or low disease activity was 40.6 percent. bDMARDs were more effective in patients with better functionality (Odds Ratio, OR = 2.140 / 95 percent Confidence Interval, CI 1.219 - 3.756) at beginning of treatment and in patients who not had a previous bDMARDs (OR = 2.150 / 95 percent CI 1.144 - 4.042).

#### **CONCLUSIONS:**

In this real-world study, functionality and use of previous bDMARDs are predictors in patients with RA treated with bDMARDs.

# OP131 Cost-Effectiveness Of Dexamethasone And Adalimumab For Uveitis

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# **INTRODUCTION:**

Uveitis is inflammation inside the eye whose underlying cause may be infectious or non-infectious. The objective of our study was to assess the cost-effectiveness of the dexamethasone implant and adalimumab compared with current practice (immunosuppressants and systemic corticosteroids) in patients with non-infectious intermediate, posterior or pan-uveitis.

# **METHODS:**

A Markov model was built to estimate costs and benefits of the interventions. Systematic reviews were performed to identify the relevant evidence. Quality of life data collected in three key randomized-controlled trials (1-3) was used to estimate the interventions effectiveness compared with the trials comparator arms, which consisted of placebo plus limited current practice (LCP). An indirect treatment comparison between adalimumab and dexamethasone was considered inappropriate due to lack of necessary evidence. For adalimumab, patients with active and inactive uveitis were considered separately. Due to the short duration of the trials, the rate of blindness, an important complication of uveitis, was highly uncertain. Substantial exploratory analyses were therefore undertaken. The analysis was performed from the perspective of the National Health Service (NHS) and