

Publication**Amiodarone-induced thyrotoxicosis: clinical course and predictors of outcome****JournalArticle (Originalarbeit in einer wissenschaftlichen Zeitschrift)****ID** 3975350**Author(s)** Conen, D.; Melly, L.; Kaufmann, C.; Bilz, S.; Ammann, P.; Schaer, B.; Sticherling, C.; Muller, B.; Osswald, S.**Author(s) at UniBasel** Müller, Beat ;**Year** 2007**Title** Amiodarone-induced thyrotoxicosis: clinical course and predictors of outcome**Journal** Journal of the American College of Cardiology**Volume** 49**Number** 24**Pages / Article-Number** 2350-5**Keywords** Aged; Amiodarone/*adverse effects; Anti-Arrhythmia Agents/*adverse effects; Diagnosis, Differential; Female; Glucocorticoids/pharmacology; Humans; Hyperthyroidism/drug therapy; Male; Middle Aged; Multivariate Analysis; Prednisone/pharmacology; Prognosis; Retrospective Studies; Thyroid Hormones/blood; Thyrotoxicosis/*chemically induced/diagnosis/drug therapy

OBJECTIVES: This study sought to determine the clinical course and predictors of long-term outcome in patients with documented amiodarone-induced thyrotoxicosis (AIT). **BACKGROUND:** Amiodarone-induced thyrotoxicosis is a condition that is difficult to manage, in particular because of the long half-life of amiodarone. Data on optimal treatment for AIT are scarce. **METHODS:** We performed a retrospective review among patients with documented AIT at a tertiary care center. Baseline characteristics, treatment received, laboratory parameters, and events during follow-up were evaluated. The predefined composite end point consisted of the following AIT-associated complications: death, heart transplantation, hospitalization for heart failure, myocardial infarction, stroke, hospitalization for arrhythmia management, or hospitalization for treatment complications. **RESULTS:** Eighty-four patients were included in the present analysis; 27 patients received prednisone for AIT. There was no difference in time to normalization of free thyroxine between those receiving and those not receiving prednisone. Long-term follow-up showed high morbidity and mortality; 47 patients (56%) reached the primary end point. Patients receiving prednisone had a worse outcome than those not receiving prednisone ($p = 0.003$). Although patients received prednisone for 84 ± 65 days, curves started to separate only 12 months after the initial diagnosis. **CONCLUSIONS:** Patients with AIT have a high event rate during follow-up. Prednisone had no effect on time to normalization of thyroxine levels and was associated with an increased event rate. Importantly, AIT-related problems must be expected late, at a time when thyroid function is under control.

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