No increased mortality in patients with rheumatoid arthritis: up to 10 years of follow up from disease onset

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Abstract

Objective—To investigate mortality, functional capacity, and prognostic factors for mortality in an inception cohort of patients with recently diagnosed RA followed up for up to 10 years.

Methods—The observed mortality of this inception cohort with recently diagnosed RA, was analysed in relation to the expected mortality, calculated with the aid of life tables of the general population of the Netherlands (matched for age and sex). Functional capacity was measured by the Health Assessment Questionnaire. Prognostic factors for mortality were analysed multivariately by the Cox proportional hazards model.

Results—Between January 1985 and April 1997, 622 patients entered the study, and were included in the analysis of mortality. The death rate in the first 10 years of the disease was not significantly different from that of the general population. Fifty five patients from the study group died (16% up to 10 years of follow up). The most commonly reported causes of death were of cardiovascular and respiratory origin. The other causes of death could be classified into cancer, sepsis, amyloidosis, leukae mia, renal insufficiency of unknown cause, perforation of the oesophagus, probably related to the treatment with non-steroidal anti-inflammatory drugs, and pancytopenia during aurothioglucose treatment. Functional capacity improved significantly during the first six years compared with the value at start. Statistically significant predictors for death were age at the start and male sex.

Conclusions—In contrast with earlier studies performed, no excess mortality in the first 10 years of an inception cohort of patients with RA was seen. In addition, the functional capacity was relatively constant during the first six years after an initial improvement from baseline. Age at start and male sex were the only statistically significant predictors for death.

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Rheumatoid arthritis (RA) is a systemic inflammatory disease characterised by chronic polyarthritis, which often leads to severe disability. Several studies have shown that RA is associated with an increased death rate.1-6 However, most of these studies show differences in their definitions of RA, selection of patients and controls, length of follow up, duration of RA before the study, methods of statistical analyses, and the linking of causes of death ascribed to RA.1-4

Only three studies investigated mortality in inception cohorts of patients with recent onset RA (symptoms for less than one year at study entry).9-12 Two studies concerned patients already included before 1975, and in both studies an increased death rate was found.9,11 In contrast, in a recently published inception cohort study no increased death rate was found, but progressive functional impairment during the first years of the disease was seen in these patients.12,13

In the past decade there has been a dramatic change in the pharmacotherapy: the number of available non-steroidal anti-inflammatory drugs (NSAIDs) has increased, but probably more importantly sulfasalazine and methotrexate, the two most commonly used disease modifying antirheumatic drugs (DMARDs) at present, were introduced in the treatment of RA. In addition, DMARDs are given earlier in the disease course, sometimes even in combinations. It has been assumed that this more aggressive treatment strategy has influenced the disease course beneficially at the long term, though this has not yet been proved.14

In several studies increased death rates have been reported for infections, especially respiratory infections, gastrointestinal related deaths, and lymphoproliferative diseases. The prevalence of cardiovascular deaths and deaths from carcinomas in patients with RA has been reported to be roughly the same as in the general population.3,15,16 We evaluated the causes of death in this present study.

In most studies the possibility of predicting mortality has also been investigated. A great variety of prognostic factors have been reported so far: age, sex, formal educational level, rheumatoid factor (RF), the presence of rheumatoid nodules at the first visit, joint count, activities of daily living, oral corticosteroid use, erythrocyte sedimentation rate (ESR), disease duration, and extra-articular manifestations of RA, particularly vasculitis.15 In the design of these studies, it is still unclear which variables are predictive for mortality.

In this study, which was started in 1985, the death rate, causes of death, functional disability, and possible prognostic factors for mortality were investigated in an inception cohort of...
patients with recent onset RA; data were collected and new patients included until 1997.

Materials and methods
Starting January 1985, all consecutive patients attending the Nijmegen or Groningen University Hospital with recent onset RA (symptoms for <1 year at study entry) according to the American Rheumatism Association criteria (ARA, 1987), were asked to participate in a long term prospective study. All patients were followed up in a standardised way and new patients were included25 up to April 1997. The patients were treated by different rheumatologists of both departments, and were seen every three months by specially trained research nurses. Quantitative clinical and laboratory data were collected, comprising the Ritchie Articular Index (RAI), the number of tender joints, number of swollen joints, ESR (according to Westergren, mm/1st h), IgM RF (enzyme linked immunosorbent assay (ELISA), normal <10 IU/ml), HLA status, and level of education. Furthermore, the patients completed a Health Assessment Questionnaire (HAQ) every six months and at each visit the disease activity score (DAS) was calculated.26 27 The starting date, dose, stopping date, reason for discontinuation, and side effects of the DMARDs and NSAIDs used were recorded.28 DMARDs included antimalarial drugs, methotrexate, sulfasalazine, gold, d-penicillamine, azathioprine, cyclophosphamide, and cyclosporin.

Standard methods of survival analysis (Kaplan-Meier, 1957) were used, in which the mortality of this RA group (n=622) was compared with the expected mortality based on the general population of the Netherlands, matched for age and sex.29 Patients lost to follow up were contacted by phone or mail with the help of family or general physicians to determine the death/life status of the relevant patient at April 1997. Causes and dates of death were obtained from the general practitioner (or rheumatologist who obtain dates and causes of death from the general practitioner in the Netherlands) and were coded according to the International Classification of Diseases, ninth revision, clinical modification (ICD-9-CM) medical diagnoses.

The functional capacity of patients attending the Nijmegen University Hospital (n=322), for whom Health Assessment Questionnaires were available at each year of follow up (n=249), are presented for each year of the follow up (table 2).

Prognostic factors for mortality were analysed multivariately by the Cox proportional hazards model, multivariate with age (start of study), sex (male or female), IgM rheumatoid factor (IgM RF positive or negative), HLA status, RAI, number of swollen joints (range 0–44), mean DAS of first year of follow up (range 0–10), and ESR fixed in the model. All variables were continuous, except sex, IgM RF, and HLA status. Patients for whom all baseline characteristics were available were included in the model (n=566). The dependent variable in the model was duration of follow up. In a separate analysis the mean HAQ of the first year was included in the Cox proportional hazards model as well.

Results
MORTALITY AND FUNCTIONAL CAPACITY
By April 1997 622 patients had entered the study and were included in the various analyses. Table 1 presents the baseline characteristics of the clinical variables of the study group. Figure 1 shows the number of patients at each time point during 10 years of follow up. The causes of death ascribed by the rheumatologists or general practitioners could be classified as cardiovascular (n=26) and respiratory cause (n=6), cancer (n=17), sepsis (n=1), amyloidosis (n=2), renal insufficiency of unknown cause (n=1), perforation of the oesophagus, probably related to NSAID treatment (n=1), and pancytopenia during aurothioglucose treatment (n=1).

Table 1  Baseline characteristics of the study group

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Patients (n=622)*</th>
</tr>
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<tbody>
<tr>
<td>Female</td>
<td>386 (62)</td>
</tr>
<tr>
<td>Age at study start (years)</td>
<td>53.3 (18–86)</td>
</tr>
<tr>
<td>IgM RF* (&gt;10 IU)</td>
<td>510 (82)</td>
</tr>
<tr>
<td>HLA-DR4+</td>
<td>336 (54)</td>
</tr>
<tr>
<td>ESR† (mm/1st hour)</td>
<td>40.7 (2.0–135.0)</td>
</tr>
<tr>
<td>Disease activity score</td>
<td>4.0 (2.8–5.6)</td>
</tr>
<tr>
<td>Ritchie Articular Index</td>
<td>12.3 (3.0–27.0)</td>
</tr>
<tr>
<td>Health Assessment Questionnaire</td>
<td>0.67 (0–1.97)</td>
</tr>
<tr>
<td>DMARD† (first year)</td>
<td>566 (91)</td>
</tr>
<tr>
<td>DMARD (follow up)</td>
<td>591 (95)</td>
</tr>
<tr>
<td>Corticosteroids (first year)</td>
<td>124 (20)</td>
</tr>
<tr>
<td>Corticosteroids (follow up)</td>
<td>212 (34)</td>
</tr>
</tbody>
</table>

*Mean values (range) and numbers (percentages) are given. †ESR = erythrocyte sedimentation rate; DMARD = disease modifying antirheumatic drug.
Confidence intervals (95%) of the study group are shown compared with the general population of the Netherlands, matched for age and sex.

Figure 2 Kaplan-Meier survival curves of patients with recent onset rheumatoid arthritis (Kaplan-Meyer) of the patients with RA (n=622) compared with the general population of the Netherlands, matched for age and sex. Confidence intervals (95%) are shown for the survival of the patients of the study group of 622 patients died with recent onset RA was analysed. Fifty five patients of the study group of 622 patients died during the period of observation. The observed mortality of this study group up to 10 years of follow up was comparable with the expected mortality, calculated with the aid of life tables, of the general population of the Netherlands (matched for age and sex). This finding is in contrast with the general belief that RA is associated with an increased death rate.

In a separate analysis, in which the mean HAQ of the first year was also included, the HAQ did not have prognostic value for mortality (results not shown).

**Discussion**

In this prospective study, started in 1985, the death rate of an inception cohort of patients with recent onset RA was analysed. Fifty five patients of the study group of 622 patients died during the period of observation. The observed mortality of this study group up to 10 years of follow up was comparable with the expected mortality, calculated with the aid of life tables, of the general population of the Netherlands (matched for age and sex). This finding is in contrast with the general belief that RA is associated with an increased death rate.

Only three studies report the mortality in patients followed up prospectively from onset. In the study of Corbett et al a small increase in mortality, compared with population expected rates, was found (these data were not analysed statistically). This excess mortality increased with increasing duration of disease, but was already apparent after three years of follow up. The study of Rasker and Cos, also including an inception cohort of patients with RA who satisfied the ARA criteria of definite or classical RA within one year of onset, did not compare their results with the general population (that is, the population of the UK). However, an increase in mortality, between the third and tenth year after baseline with a loss of life expectancy of at least five years was suggested. A striking difference between the study of Rasker and Cos and our data is the mean age at death (73 years in this study v 68 years in the study of Rasker et al). In contrast, only the study of Lindquist and Eberhardt found no increased mortality.

The causes of death observed in our study were comparable with the results seen in the three inception cohort studies, except for two deaths related to treatment which occurred in this study. In the three previous studies second line drugs were not responsible for any...
In the study of Lindquist and Eberhardt only 62% of the patients were treated with DMARDs and 16% with oral corticosteroids at some time during follow up. In our study 91% of the patients were already being treated with DMARDs during the first year of follow up and 95% of the patients were treated with DMARDs at some time during follow up. In addition, 20% of the patients used oral corticosteroids at some time during the first year of observation, and 35% of the patients used oral corticosteroids at some time during the complete follow up period. In the Netherlands there is a general policy of referring patients with RA early in the disease course to a rheumatologist. Therefore none of the patients had started taking DMARDs or corticosteroids before inclusion in the study. Although it seems logical to suggest that the early use of second line treatment might lead to treatment related mortality, this was not confirmed by our study as no increased mortality was found. In other mortality studies increased death rates in patients with RA were reported as a result of infection, especially respiratory infection, gastrointestinal and lymphoproliferative diseases. However, most of these studies had different definitions of RA, disease duration before study inclusion, and length of follow up, and most patients received steroid treatment for several years. The functional capacity was relatively constant during the first six years after an initial improvement from baseline. After 10 years of follow up the functional capacity had returned to the level at the start of the study. This is in contrast with the study of Corbett et al., in which disability was reported to develop most rapidly during the first year after disease onset, with a slow, nearly linear rate of increase during the period of observation. In the study of Eberhardt and Fex, in which no increased mortality was found in an inception cohort of patients with RA, it was concluded that functional impairment of different joints had progressed, but most patients were mildly disabled up to six years of follow up. In that study fewer patients were treated with second line drugs or corticosteroids during follow up, and therefore we suggest that the early use of second line treatment in our study, in almost all patients, might have led to improved functional capacity but not to increased mortality.

Different variables at onset were also analysed for their prognostic value for mortality. No univariate analysis was performed, as mortality and various disease activity variables are highly age and sex related. Therefore age would act as a confounder in a univariate analysis. In the regression model only age at start and male sex were statistically significant prognostic factors. However, Anderson, who reviewed 25 articles on age and mortality in RA, failed to show a clear association between sex, age, and mortality in RA.

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5 Gabriel SE, Crowson CS, O’Fallon WM. Mortality in rheumatoid arthritis: have we made an impact in 4 decades? J Rheumatol 1999;26:2529–33.


