Evidence for differential acquired drug resistance to anti-tumour necrosis factor agents in rheumatoid arthritis

A Finckh, J F Simard, C Gabay, P-A Guerne, for the SCQM physicians

Background: Acquired drug resistance or gradual drug failure has been described with most disease modifying antirheumatic drugs (DMARDs) and is also starting to be recognised with anti-tumour necrosis factor (anti-TNF) agents.

Objective: To study acquired drug resistance to anti-TNF agents in rheumatoid arthritis (RA).

Methods: Swiss health authorities requested continuous monitoring of patients receiving biological agents. Intensification of co-therapy with traditional DMARDs, gradual dose escalation, and drug discontinuation rates in all patients receiving infliximab, etanercept, or adalimumab, adjusting for potential confounders, were analysed. Intensification of DMARD co-therapy and time to discontinuation of the three anti-TNF agents were analysed using a proportional hazards models. Dose escalation and evolution of RA disease activity (DAS28) were analysed using a longitudinal regression model.

Results: 1198 patients contributing 1450 patient-years of anti-TNF treatment met the inclusion criteria. The rate of intensification of traditional DMARD co-therapy over time was significantly higher with infliximab (hazards ratio = 1.73 (99% confidence interval (CI) 1.19 to 2.51)) than with the two other agents.

Infliximab also showed significant dose escalation over time, with an average dose increase of +12% (99% CI 8% to 16%) after 1 year, and +18% (99% CI 11% to 25%) after 2 years. No significant differences in discontinuation rates were seen between the three anti-TNF agents (ANOVA, p = 0.67). Evolution of disease activity over time indicated a lower therapeutic response to infliximab (DAS28, p < 0.001) compared with etanercept, after 6 months' treatment.

Conclusions: In this population, infliximab was associated with a higher risk of requiring intensification of DMARD co-therapy than the other anti-TNF agents and a significant dose escalation over time. Analysis of RA disease activity indicated a reduced therapeutic response to infliximab after the first 6 months of treatment, suggestive of acquired drug resistance.

New disease modifying antirheumatic drugs (DMARDs) have become available since 1999. Biological agents, such as anti-tumour necrosis factor α (anti-TNF), dramatically improve the signs and symptoms of rheumatoid arthritis (RA) refractory to conventional treatment. The chemical structure, pharmacokinetic properties, and specific mechanisms of TNF inhibition of available anti-TNF agents differ: infliximab (INF) (Remicade; Centocor INC, Malvern, PA, USA) is a chimeric monoclonal anti-TNF antibody (human IgGk/mouse Fc); adalimumab (ADL) (Humira; Abbott Laboratories, Illinois, USA) is a fully human monoclonal anti-TNF antibody; whereas etanercept (ETN) (Enbrel; Amgen, Inc, Thousand Oaks, CA, USA) is an engineered TNF receptor (humanised protein) acting as a competitive inhibitor of TNFα and β. The unique pharmacological properties of these agents have been associated with different rates of opportunistic granulomatous infections and are thought to explain why some anti-TNF agents work in some chronic inflammatory conditions and not in others. It is not known whether these differences affect their long term therapeutic effectiveness or the potential development of drug resistance in RA.

In clinical practice, loss of effectiveness of long term DMARD treatment is a common problem. Acquired drug resistance or gradual drug failure has been described with most traditional DMARDs and is also starting to be recognised with anti-TNF agents. Not all patients with RA respond to the standard dosage of anti-TNF agents; 28–58% of all patients with RA show little response to these drugs in large randomised trials. Acquired resistance to DMARD treatment in RA has been measured by analysing use of additional DMARD co-therapy, anti-TNF dose escalation, and drug discontinuation rates (“drug survival”). These outcomes reflect the common therapeutic options a physician has when faced with loss of DMARD effectiveness: for patients not fully responsive to anti-TNF agents, physicians may increase co-therapy with traditional DMARDs, increase the anti-TNF dose, or decide to stop the current anti-TNF treatment and switch to other treatments. Dosage escalation has been observed with infliximab, but this may not be a valid measure of drug resistance for anti-TNF agents without a flexible dosing regimen (ETN, ADL), in which case intensification of traditional DMARD co-therapy and drug discontinuation might be more adequate outcomes.

This study aimed at investigating acquired drug resistance to anti-TNF treatments in a population based observational cohort of patients with RA. We examined intensification of DMARD co-therapy, progressive dose escalation, and drug discontinuation rates in all patients receiving infliximab, etanercept, or adalimumab, adjusting for potential confounders, were analysed. Intensification of DMARD co-therapy and time to discontinuation of the three anti-TNF agents were analysed using a proportional hazards models. Dose escalation and evolution of RA disease activity (DAS28) were analysed using a longitudinal regression model.

Conclusions: In this population, infliximab was associated with a higher risk of requiring intensification of DMARD co-therapy than the other anti-TNF agents and a significant dose escalation over time. Analysis of RA disease activity indicated a reduced therapeutic response to infliximab after the first 6 months of treatment, suggestive of acquired drug resistance.

Abbreviations: ADL, adalimumab; ANOVA, analysis of variance; CI, confidence interval; DAS28, 28 joint count Disease Activity Score; DMARDs, disease modifying antirheumatic drugs; ETN, etanercept; HAQ, Health Assessment Questionnaire; HR, hazard ratio; INF, infliximab; RA, rheumatoid arthritis; RF, rheumatoid factor; SCQM, Swiss Clinical Quality Management; TNF, tumour necrosis factor;
discontinuation rates of the three available anti-TNF agents. In addition, we explored underlying pathways leading to these therapeutic adjustments in relationship to RA disease activity.

**PATIENTS AND METHODS**

**Study population**

Regulatory agencies in Switzerland have requested continuous monitoring of all patients receiving costly biological agents. The Swiss Clinical Quality Management of RA (SCQM) system was established by the Swiss Society of Rheumatology and elected to follow up all patients with RA starting to receive anti-TNF agents. The patient’s rheumatologist or primary care physician are incited to enrol their patient in the SCQM by allowing them to deduct the costs of anti-TNF drugs from their global treatment expenditure scrutinised by the health authorities; this contributes to a high enrolment rate. Based on a comparison with sales data from the industry, between 70 and 80% of all Swiss patients with RA receiving anti-TNF agents are included in the SCQM. Patients are enrolled at the start of anti-TNF therapy and followed up prospectively. The SCQM includes measurements of disease activity, radiographic damage, adverse drug reactions, and RA symptoms. Clinical information is collected systematically by the patient’s physician every 6–12 months and further updated at every significant change in antirheumatic treatment. The accuracy of medication data provided by the physicians—including start and stop dates—was confirmed against records from the pharmaceutical industry and participants’ self reported information. Patients come from a wide variety of clinical settings: 40% private rheumatology practices, 30% non-academic centres, and 30% academic centres. This analysis includes data collected between January 1998 and the end of September 2004. The inclusion criteria for this analysis were a diagnosis of RA by a rheumatologist and treatment with INF, ETN, or ADL.

**Outcomes**

We considered three outcomes that operationally define drug resistance: increase in concomitant DMARD treatment, dosage escalation in anti-TNF agents, and interruption of the current anti-TNF therapy. Traditional DMARD co-therapy was defined as concomitant prescription of methotrexate, leflunomide, sulfasalazine, ciclosporin, hydroxychloroquine, or azathioprine with an anti-TNF agent. Increase in concomitant DMARD treatment was defined by a rise in the weekly dose of traditional DMARDs or by the introduction of a new DMARD in addition to the current treatment. Replacement of one concomitant DMARD by another was not considered an increase in DMARD co-therapy. Dose escalation can occur either by increasing the dose of an anti-TNF agent or by shortening the dispensation interval. To take into account both of these possibilities, we computed each patient’s average weekly dose of anti-TNF therapy (mg/week) at each time point. We examined drug retention of anti-TNF agents started to become statistically different only after 12 months and further updated at every significant change in antirheumatic treatment. The accuracy of medication data provided by the physicians—including start and stop dates—was confirmed against records from the pharmaceutical industry and participants’ self reported information. Patients come from a wide variety of clinical settings: 40% private rheumatology practices, 30% non-academic centres, and 30% academic centres. This analysis includes data collected between January 1998 and the end of September 2004. The inclusion criteria for this analysis were a diagnosis of RA by a rheumatologist and treatment with INF, ETN, or ADL.

**RESULTS**

A total of 1198 patients receiving anti-TNF treatment met the inclusion criteria, with assessments every 4 months on average. Some differences in baseline characteristics between the three groups were noted (table 1). In particular, for those receiving ADL another anti-TNF agent had failed in a higher proportion (p=0.006), and they had slightly lower disease activity, and less functional disability (HAQ) at baseline.

**Intensification of DMARD co-therapy**

The rate of intensification of traditional DMARD co-therapy was significantly different between the three treatments (ANOVA, p = 0.006) (fig 1). The hazard ratio (HR) of increasing traditional DMARD co-therapy over time with INF was 1.73 (99% CI 1.19 to 2.51) compared with the other anti-TNF agents, while there was no significant difference between ADL and ETN (Bonferroni’s p = 0.82). The difference in DMARD co-therapy intensification between anti-TNF agents started to become statistically different only after
Table 1  Baseline characteristics of patients at initiation of anti-TNF therapy

<table>
<thead>
<tr>
<th>Disease characteristics*</th>
<th>Adalimumab (ADL) (n = 317)</th>
<th>Infliximab (INF) (n = 362)</th>
<th>Etanercept (ETN) (n = 519)</th>
<th>p Value†</th>
</tr>
</thead>
<tbody>
<tr>
<td>Anti-TNF dose, median (IQR)</td>
<td>40 mg/2 wk (40–40)</td>
<td>3.2 mg/kg x 8 wk (3.0–3.75)</td>
<td>50 mg/wk (50–50)</td>
<td></td>
</tr>
<tr>
<td>Female (%)</td>
<td>74</td>
<td>75</td>
<td>74</td>
<td>0.89</td>
</tr>
<tr>
<td>Age (years)</td>
<td>53 (51 to 54.7)</td>
<td>53.1 (51.7 to 54.5)</td>
<td>54.4 (53.2 to 55.6)</td>
<td>0.24</td>
</tr>
<tr>
<td>Disease duration (years), median (IQR)</td>
<td>10.1 (5.6–17.5)</td>
<td>10.2 (5.0–16.5)</td>
<td>10.3 (5.7–15.9)</td>
<td>0.97</td>
</tr>
<tr>
<td>Rheumatoid factor+ (%)</td>
<td>82</td>
<td>81</td>
<td>78</td>
<td>0.33</td>
</tr>
<tr>
<td>Follow up (mo; median (IQR)</td>
<td>10.7 (5.8–12.3)</td>
<td>18.8 (11.5–28.3)</td>
<td>23.7 (12.6–35.8)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Previous failure of anti-TNF agent (%)</td>
<td>39</td>
<td>37</td>
<td>37</td>
<td>0.98</td>
</tr>
<tr>
<td>Disease activity score (DAS28)</td>
<td>4.19 (4.02 to 4.36)</td>
<td>4.54 (4.38 to 4.7)</td>
<td>4.72 (4.59 to 4.85)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>RADAI</td>
<td>4.02 (3.6 to 4.5)</td>
<td>4.59 (4.37 to 4.81)</td>
<td>4.85 (4.67 to 5.03)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Function disability (HAQ)</td>
<td>1.25 (1.18 to 1.33)</td>
<td>1.37 (1.29 to 1.44)</td>
<td>1.37 (1.31 to 1.43)</td>
<td>0.04</td>
</tr>
<tr>
<td>Concomitant DMARD use (%)</td>
<td>53</td>
<td>93</td>
<td>64</td>
<td></td>
</tr>
<tr>
<td>Methotrexate (%)</td>
<td>37</td>
<td>70</td>
<td>40</td>
<td></td>
</tr>
<tr>
<td>Leflunomide (%)</td>
<td>10</td>
<td>10</td>
<td>10</td>
<td></td>
</tr>
<tr>
<td>Sulfasalazine (%)</td>
<td>4</td>
<td>6</td>
<td>6</td>
<td></td>
</tr>
<tr>
<td>Other DMARD (%)</td>
<td>2</td>
<td>7</td>
<td>8</td>
<td></td>
</tr>
<tr>
<td>Glucocorticoid use (%)</td>
<td>41</td>
<td>56</td>
<td>60</td>
<td>&lt;0.001</td>
</tr>
</tbody>
</table>

*Values are given as means with 95% confidence intervals (CI) if not otherwise indicated. When not normally distributed variables, medians and interquartile ranges (IQRs) are reported; + Test for dichotomous variables; † ANOVA, p = 0.006.

1.2 years of treatment. Significant predictors of DMARD intensification were previous failure of another anti-TNF agent (HR = 2.14, 95% CI 1.42 to 3.23), RF positivity (HR = 1.64, 95% CI 1.07 to 2.51), and disease duration (HR = 0.98, 95% CI 0.96 to 0.99). As expected, starting INF without concomitant DMARDs significantly increased the need for subsequent intensification of co-therapy (HR = 4.51, 95% CI 2.64 to 7.71), but this was not the case with ETN or ADL. No difference was observed between anti-TNF agents in glucocorticoid intensification or glucocorticoid reduction. Various sensitivity analyses with different definitions of DMARD co-therapy intensification, accounting for simultaneous changes in glucocorticoid dose, provided very similar results.

**Dose escalation**

Dose escalation was also significantly different between the three anti-TNF agents (ANOVA, p < 0.001), with only INF demonstrating a significant change in dosage over time (fig 2). The average dose escalation was 3.01 mg/week (99% CI 2.0 to 4.03) after the first year of INF treatment, which represents an average increase of 24 mg per 2 month period or +12% in INF dose. Furthermore, the average INF dose continued to escalate significantly after the first year of treatment and reached 4.60 mg/week (99% CI 2.84 to 6.36), which represents an average increase of 37 mg per 2 month period or +18% in INF dose. Dose escalation of INF was significantly higher in patients without concomitant DMARD treatment (5.5 mg/week at 1 year, 99% CI 1.9 to 9.01). No significant change in dose from baseline was apparent with ETN (Bonferroni, p = 0.08) or with ADL (Bonferroni,
Improvement of 0.13 (99% CI 0.03 to 0.27) with ETN during the following year, compared with a non-significant INF the DAS28 worsened by 0.14 (99% CI 0.01 to 0.27) across the three groups, but thereafter the curves separated: with rates of improvement in disease activity varied little between anti-TNF agents with concomitant DMARDs, discontinuation therapy on anti-TNF discontinuation: in patients starting anti-TNF agent increased the rate of drug discontinuation by 77% (HR = 1.77, 95% CI 1.23 to 2.54) in a crude analysis. In a subgroup analysis, we explored the effect of DMARD co-therapy on anti-TNF discontinuation: in patients starting anti-TNF agents with concomitant DMARDs, discontinuation rates were significantly lower with ETN (HR = 0.66, 99% CI 0.45 to 0.98) than with INF or ADL.

**Therapeutic effectiveness on RA disease activity**

The evolution of RA disease activity (DAS28) over time differed significantly between anti-TNF agents (ANOVA, p = 0.06). However, progressive dose escalation may not be a valid measure of drug resistance for anti-TNF agents not commonly used with a flexible dosing regimen (ETN, ADL). Of the other covariates, only RF positivity significantly influenced changes in dose (p = 0.025).

**Discontinuation of treatment**

The discontinuation rates were not significantly different between the treatments (ANOVA, p = 0.67) (fig 3). A total of 1450.5 patient-years of anti-TNF therapy and 300 cases of treatment interruption were examined. The overall discontinuation rate of anti-TNF agents was relatively low with a median drug survival of 3.21 years (IQR = 1.43–4.52). The HR of discontinuing treatment was 1.11 (99% CI 0.89 to 1.40) for INF compared with ETN and ADL, 0.97 (99% CI 0.71 to 1.15) for ADL compared with INF and ETN, and 0.91 (99% CI 0.71 to 1.15) for ETN compared with INF and ADL. Significant predictors of discontinuation were increased disability at baseline as measured by the HAQ (HR = 1.22, 95% CI 1.00 to 1.48) and calendar year of treatment initiation (HR = 2.01, 95% CI 1.71 to 2.36), which reflects the increasing proportion of patients starting biological agents after previous anti-TNF agents have failed and a greater availability of therapeutic alternatives favouring treatment switches over time. As expected, previous failure of another anti-TNF agent increased the rate of drug discontinuation by 77% (HR = 1.77, 95% CI 1.23 to 2.54) in a crude analysis. In a subgroup analysis, we explored the effect of DMARD co-therapy on anti-TNF discontinuation: in patients starting anti-TNF agents with concomitant DMARDs, discontinuation rates were significantly lower with ETN (HR = 0.66, 99% CI 0.45 to 0.98) than with INF or ADL.

**DISCUSSION**

We studied operational features of drug resistance to anti-TNF agents in RA by analysing intensification of DMARD co-therapy, gradual dose escalation, and drug discontinuation rates in a population based cohort of 1196 patients with RA receiving INF, ADL, or ETN. Intensification of traditional DMARD co-therapy over time was significantly higher with INF than with the other anti-TNF agents and progressive dose escalation was evident for INF, but not for ETN or ADL. No significant difference in discontinuation rates was seen. Analyses of RA disease activity over time indicated a lower rate of intensification of DMARD co-therapy after previous anti-TNF agents have failed and a greater availability of therapeutic alternatives favouring treatment switches over time. As expected, previous failure of another anti-TNF agent increased the rate of drug discontinuation by 77% (HR = 1.77, 95% CI 1.23 to 2.54) in a crude analysis. In a subgroup analysis, we explored the effect of DMARD co-therapy on anti-TNF discontinuation: in patients starting anti-TNF agents with concomitant DMARDs, discontinuation rates were significantly lower with ETN (HR = 0.66, 99% CI 0.45 to 0.98) than with INF or ADL.

**Therapeutic effectiveness on RA disease activity**

The evolution of RA disease activity (DAS28) over time differed significantly between anti-TNF agents (ANOVA, p = 0.001) (fig 4). During the first 6 months of treatment, rates of improvement in disease activity varied little between the three groups, but thereafter the curves separated: with INF the DAS28 worsened by 0.14 (99% CI 0.01 to 0.27) during the following year, compared with a non-significant improvement of 0.13 (99% CI 0.03 to 0.27) with ETN during the same period. Data for ADL were insufficient beyond 1 year of treatment to allow reliable comparisons. After the first year of treatment, the DAS28 had improved by 0.68 (99% CI 0.30 to 1.07) with ADL, by 0.82 (99% CI 0.61 to 1.03) with INF, and by 1.05 (99% CI 0.87 to 1.23) with ETN. Depending on the absolute level of disease activity, DAS28 improvements between 0.6 and 1.2 are considered good therapeutic responses and improvements >1.2 are considered good therapeutic responses (EULAR response criteria). After the first year of treatment, 47% (99% CI 35% to 59%) of patients were classified as having a moderate or good response to ADL (99% CI 44% to 61%) to INF, and 66% (99% CI 59% to 72%) to ETN.
Not all patients with RA respond to the suggested initial INF dose of 3 mg/kg. Higher serum levels of INF in the ATTRACT trial were associated with increased clinical response, reduction in C reactive protein levels, and reduced radiographic joint damage progression, which suggested a dose-response relationship.\(^5\) Subsequently, a flexible dosing regimen has been accepted for INF in some countries, including Switzerland, which allows dose increases up to 10 mg/kg and reductions in the dosing interval to every 4 weeks. Gradual dose escalation of INF has since been observed in several RA cohorts.\(^{20-25}\) The dose increase in an American cohort was 36% at 1 year,\(^{26}\) significantly more than in our cohort (12%), which might reflect a difference in clinical practice and in patient population. It is not clear whether this dose escalation reflects initial dose adjustments of INF therapy or acquired drug resistance to this agent. In this study and in others,\(^{27}\) dose escalation of INF was steepest during the first year of treatment, but continued to increase in the second year at a slower rate, suggesting that both phenomena may be involved. Because of the price of anti-TNF therapy, these results have important economic and clinical implications. It has been estimated that INF therapy incurred a 25% increase in 1-year costs representing an average of $4200 per patient per year in the USA.\(^{28}\) A higher INF dosage may also carry an increased risk of side effects.\(^{20-27}\)

No significant difference in discontinuation rates was seen between the three anti-TNF agents (p = 0.67) after adjusting for confounders. This corroborates results of other studies on retention rates of anti-TNF agents.\(^{38-40}\) Drug discontinuation rates represent a useful summary measure of overall effectiveness of a treatment, even if the rate is influenced by the availability of therapeutic alternatives or the incidence of adverse drug reactions. Anti-TNF agents had relatively low discontinuation rates and median drug survival of 3.21 years (IQR = 1.43–4.52) in this population. Severe adverse drug reactions were relatively infrequent in this cohort,\(^{31}\) suggesting that drug discontinuation denoted mostly unsatisfactory treatment response.\(^{31}\)

In usual clinical practice, concomitant DMARD co-therapy or anti-TNF agents are increased only if disease activity is not satisfactorily controlled, and tapered only when disease activity is adequately reduced. To link the therapeutic adjustments with clinically important outcomes, we examined RA disease activity over time in the three treatment groups using an independent outcome measure. The average effect size of the DAS28 response to anti-TNF agents was moderate, with an overall improvement of 0.92 (95% CI 0.81 to 1.03) after the first year.\(^{44}\) Similar values of DAS28 improvement have been observed in other cohorts: in Swedish patients with RA, INF treatment was associated with a median DAS28 improvement of 0.6 after 1 year.\(^{45}\) However, the course of disease activity differed significantly between INF and the other anti-TNF agents. After a similar initial improvement, RA disease activity tended to slow or increase with INF after the first 6 months of treatment compared with ETN. Smaller therapeutic responses on RA disease activity were associated with dose escalation and concomitant DMARD treatment intensification in INF treated patients. While the mechanisms of resistance to INF or other anti-TNF agents are still poorly understood, the presence of antibodies to INF has been shown to reduce clinical response to this drug in patients with Crohn’s disease\(^{46-48}\) and is also thought to contribute to the loss of response over time in some patients with RA. Unique pharmacological properties of anti-TNF agents have been associated also with differences in the incidence of opportunistic granulomatous infections\(^{5-7}\) or the efficacy profiles in other chronic inflammatory conditions.\(^{7-9}\) Further research is needed to understand the mechanisms leading to loss of drug effectiveness with anti-TNF agents.

Some potential limitations inherent to the analysis of observational data need discussion. Firstly, in this study, there was no control over the treatment assignment. Because no rationale exists for considering that one anti-TNF agent is more efficacious than another,\(^7\) major confounding by indication between these agents is unlikely. However, ADL had a higher proportion of patients for whom another anti-TNF agent had failed, slightly lower disease activity, and less functional disability (HAQ) at baseline. These differences probably reflect the fact that ADL is the latest of the three anti-TNF agents to come on the market and that indication for these drugs has widened with time. While we were able to adjust our analysis for potential confounding by levels of disease activity, functional disability, RF positivity, previous failure of anti-TNF agents, or time trends, we could not exclude the possibility of confounding by unmeasured factors. For example, we could not adjust for concomitant rheumatic diseases or for self medication use such as analgesics. Although potential unmeasured confounding is always a concern, we have no evidence of a systematic bias, which would imply a differential prevalence of comorbidities or of co-medication use between anti-TNF agents. Intensification or reduction of concomitant glucocorticoid treatment was not significantly different between anti-TNF agents.

Secondly, missing data are another concern with observational studies. To deal with this, we checked the accuracy of medication data provided by the physicians—including start and stop dates—against records from the pharmaceutical industry and participants’ self report. However, some residual non-differential misclassification is possible, which would tend to bias the results towards the null.\(^{43}\) We included all patients with RA receiving anti-TNF agents since 1999, but for some patients (31%, table 1) no follow up data were available yet, allowing these patients to contribute only baseline information. The median follow up of patients receiving ADL, in particular, was only 10.7 (IQR = 5.8–12.3) months because it is the last agent to be introduced, which limited the ability to examine its potential drug resistance. By far, the most common reason for incomplete follow up was the recent initiation of anti-TNF therapy, with insufficient time for follow up measurements. Baseline disease characteristics of patients without complete follow up were similar to those included in the present analysis (data not shown), suggesting that the subjects in the analysis were a representative sample of the whole population.

The definition of acquired therapeutic resistance for antirheumatic treatment is less characterised than for antimicrobial treatment, for example.\(^{49}\) Acquired resistance to DMARD treatment has been measured by analysing the use of additional DMARD co-therapy,\(^{50}\) anti-TNF dose escalation,\(^{11}\) and drug discontinuation rates,\(^{8-10,16-18}\) which are only indirect measures of this concept. Further research is needed to understand the mechanisms of drug resistance to DMARDs and define outcome measures of drug resistance in RA. Moreover, we could not adjust this study’s longitudinal analyses for changes in time dependent confounders such as simultaneous variations in dose of glucocorticoid or DMARD co-therapy, because this would have introduced biases.\(^{49}\) We accounted for changes in dose of glucocorticoid and DMARD co-therapy by examining intensification or reduction of these
treatments over time independently. Strengths of this analysis include a true population based cohort, as all patients with RA receiving anti-TNF agents are requested to be enrolled by the Swiss authorities, and a systematic prospective ascertainment of a wide variety of disease characteristics.

In this population, INF was associated with a higher risk of requiring intensification of DMARD co-therapy and a significant dose escalation compared with the other anti-TNF agents. Analyses of the evolution of RA disease activity indicated a reduced therapeutic response to INF after the first 6 months of treatment, suggesting gradual drug resistance to this treatment. Further research is needed to study the mechanisms of drug resistance to INF and other anti-TNF agents.

ACKNOWLEDGEMENTS

Axel Finckh is supported by the Kirkland Scholars Fellowship, by the Centocor Health Outcomes in Rheumatic Diseases (CHORD) fellowship, by an unrestricted research grant from Wyeth, and by NIH grant P60 AR47782. Julia F Simard is supported by the Kirkland Scholars Fellowship and by NIH grant T32 AI071535. We are grateful to the SCQM staff for data management and support and to participating physicians and patients who made this study possible, to Dr Liang and Dr Costenbader, BWH, Boston for critical input into the manuscript.

SCQM has received grants from the Swiss Health authorities (BAG), the Swiss Academy for Medical Sciences (SAMW), and private companies (Wyeth, Abbott, Essex, Aventis, Bristol-Mayers, Mepha, Merck, Novartis, and Roche).

This work was supported by the SCQM staff for data management and support, with the Swiss Health authorities, and a systematic prospective ascertainment of a wide variety of disease characteristics. Strengths of this analysis include a true population based cohort, as all patients with RA receiving anti-TNF agents are requested to be enrolled by the Swiss authorities, and a systematic prospective ascertainment of a wide variety of disease characteristics.
Evidence for differential acquired drug resistance to anti-tumour necrosis factor agents in rheumatoid arthritis
A Finckh, J F Simard, C Gabay and P-A Guerne

Ann Rheum Dis 2006 65: 746-752 originally published online December 8, 2005
doi: 10.1136/ard.2005.045062

Updated information and services can be found at:
http://ard.bmj.com/content/65/6/746

These include:

References
This article cites 41 articles, 19 of which you can access for free at:
http://ard.bmj.com/content/65/6/746#BIBL

Email alerting service
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Topic Collections
Articles on similar topics can be found in the following collections

- Connective tissue disease (4253)
- Degenerative joint disease (4641)
- Immunology (including allergy) (5144)
- Musculoskeletal syndromes (4951)
- Rheumatoid arthritis (3258)
- Biological agents (545)
- Drugs: musculoskeletal and joint diseases (700)

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/