Influence of TNF-α inhibition on oxidative stress of rheumatoid arthritis patients

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SUMMARY
The aim of this study was to assess circulating levels of reactive oxygen metabolites (ROMs) as a marker of oxidative stress in rheumatoid arthritis (RA) patients during an anti-tumor necrosis factor alpha (TNF-α) treatment.

We enrolled 40 patients with RA (36 females; age 53±13 yrs) treated with different subcutaneously administered TNF-α inhibitors. The oxidative status was determined on the basis of plasma samples taken before, at 24 and 52 weeks of the anti-TNF-α treatment. Hydroperoxide levels were measured using the d-ROMs test, a useful clinically proven oxidative stress marker.

During the anti-TNF-α therapy, we observed a significant reduction in serum ROMs levels in RA patients from 33.2±10 mg H2O2/L at baseline to 29.5±7 and 29.3±9 mg H2O2/L, at 24 and 52 weeks, respectively (p<0.05). We also identified a significant correlation between the oxidative stress status and the disease activity score on 28 joints/C-reactive protein and health assessment questionnaire disability index. The results of our study demonstrate that a good control of the disease with anti-TNF-α agents can reduce oxidative stress in RA patients. However, further studies of larger patient cohorts are needed to confirm these preliminary data.

Key words: Anti-TNF; oxidative stress; rheumatoid arthritis; reactive oxygen species.

INTRODUCTION
Rheumatoid arthritis (RA) is a chronic, systemic inflammatory disease that affects primarily the joints with synovial hyperplasia and may result in structural joint damage and disability (1). The combination of genetic susceptibility along with various environmental factors is reported to play a critical role in the pathogenesis of this disease and leads to persistent immune system activation and highly complex and coordinated processes that initiate and perpetuate an autoimmune response with a low-grade chronic inflammation (2).

Reactive oxygen species (ROS) are chemically reactive molecules containing oxygen generated during normal oxygen metabolism, which play important roles in cell signaling and homeostasis. However, an imbalance between the systemic production of ROS and the biological system’s ability to readily detoxify these metabolites results in oxidative stress in cells with an ensuing damage to their components, such as proteins, lipids, and DNA (3).

ROS have been classically described as harmful products released during inflammatory diseases that can activate and perpetuate the immune response (4). In several autoimmune diseases, pro-inflammatory compounds (cytokines and prostaglandins) together with ROS are released at sites of inflammation. In particular, the overproduction of tumor necrosis factor alpha (TNF-α) is thought to be the main contributor to increased ROS release in patients with RA (5). Therefore, high ROS levels have proven to be related to disease activity (6).

The aim of this study was to assess the effect of an anti-TNF-α treatment on circu-
lating levels of reactive oxygen metabolites (ROMs) and oxidative stress in patients with active RA.

**PATIENTS AND METHODS**

According to the 2010 classification criteria of the American College of Rheumatology/European League Against Rheumatism (ACR/EULAR) (7), we consecutively enrolled 40 early RA patients, who failed treatment with disease modifying antirheumatic drugs (DMARDs) and were eligible for a biologic therapy, in compliance with the recommendations of the Italian Society of Rheumatology (8). All patients were evaluated for disease activity by the disease activity score on 28 joints/C-reactive protein (DAS28-CRP), functional status by the health assessment questionnaire disability index (HAQ-DI), and were screened for latent tuberculosis, B and C hepatitis as well as human immunodeficiency virus and varicella zoster virus exposure/infection, and history of previous malignancies. Smokers of more than 10 cigarettes/day were excluded. The clinical features of the enrolled patients are reported in Table I. Corticosteroids (CCS) and/or DMARDs treatment at baseline were

**Table I - Clinical features of rheumatoid arthritis patients (N. 40).**

<table>
<thead>
<tr>
<th></th>
<th>Baseline</th>
<th>24 weeks</th>
<th>52 weeks</th>
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</thead>
<tbody>
<tr>
<td>Female, n. (%)</td>
<td>36 (90%)</td>
<td></td>
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<tr>
<td>Age, years - mean±SD</td>
<td>53±13</td>
<td></td>
<td></td>
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<tr>
<td>Disease duration, months - mean±SD</td>
<td>6.8±3.7</td>
<td></td>
<td></td>
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<tr>
<td>RF positive, n. (%)</td>
<td>31 (77.5)</td>
<td></td>
<td></td>
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<tr>
<td>ACPA positive, n. (%)</td>
<td>29 (72.5)</td>
<td></td>
<td></td>
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<tr>
<td>Treatment, n. (%)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>CCS</td>
<td>3 (75)</td>
<td>27 (67.5)</td>
<td></td>
</tr>
<tr>
<td>CCS+MTX</td>
<td>4 (10.0)</td>
<td>10 (25.0)</td>
<td></td>
</tr>
<tr>
<td>CCS+LFM</td>
<td>6 (15.0)</td>
<td></td>
<td></td>
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<tr>
<td>CCS+SSZ</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Anti-TNF</td>
<td>15 (37.5%)</td>
<td></td>
<td></td>
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<tr>
<td>Etanercept</td>
<td>17 (42.5%)</td>
<td></td>
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<tr>
<td>Adalimumab</td>
<td>8 (20%)</td>
<td></td>
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<tr>
<td>Golimumab</td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>ESR, mm/h - mean±SD</td>
<td>57±27</td>
<td>22±21*</td>
<td>18±15*</td>
</tr>
<tr>
<td>CRP, mg/L - mean±SD</td>
<td>6.4±3.7</td>
<td>2.1±1.4*</td>
<td>1.3±1.2*</td>
</tr>
<tr>
<td>DAS28-CRP - mean±SD</td>
<td>6.4±0.9</td>
<td>3.5±1.2*</td>
<td>3.3±1.1*</td>
</tr>
<tr>
<td>EULAR response: none/moderate/good - n. (%)</td>
<td>-</td>
<td>4/14/22**</td>
<td>0/9/27**</td>
</tr>
<tr>
<td>[DAS28 improvement &lt;0.6/0.6-1.2/&gt;1.2]</td>
<td>-</td>
<td>(10/35/55)</td>
<td>(0/25/75)</td>
</tr>
<tr>
<td>HAQ-DI - mean±SD</td>
<td>1.8±0.7</td>
<td>0.9±0.4*</td>
<td>0.7±0.5*</td>
</tr>
<tr>
<td>ROS serum levels - mean±SD (mgH$_2$O$_2$/L)</td>
<td>33.2±10</td>
<td>29.5±9*</td>
<td>29.3±9*</td>
</tr>
<tr>
<td>Oxidative stress: low/moderate/high - n. (%)</td>
<td>0/17/23</td>
<td>20/16/4**</td>
<td>28/8/0**</td>
</tr>
<tr>
<td>[ROS levels &lt;27/27-32/&gt;32 mgH$_2$O$_2$/L]</td>
<td>(0/43/57)</td>
<td>(50/40/10)</td>
<td>(78/22/0)</td>
</tr>
</tbody>
</table>

SD, standard deviation; RF, rheumatoid factor; ACPA, anti-citrullinated peptide antibodies; CCS, corticosteroids; MTX, methotrexate; LFM, luflunomide; SSZ, sulfasalazine; TNF, tumor necrosis factor; ESR, erythrosedimentation rate; CRP, C-reactive protein; DAS28, disease activity score on 28 joints; EULAR, European League Against Rheumatism; HAQ-DI, health assessment questionnaire disability index; ROS, reactive oxygen species. *p<0.05; **p<0.001.
maintained in combination with TNF-α inhibitors and dosages remained unchanged during the study. At baseline and after 24 and 52 weeks from the initiation of treatment with anti-TNF-α drugs for all patients, the four-variable DAS28-CRP was measured and blood samples were taken and frozen at -80°C until laboratory assessment. Plasma levels of ROMs were measured by performing the Diacron reactive oxygen metabolite d-ROM test, according to the manufacturer’s instructions. This test reflects the amount of organic hydroperoxides that is related to the free radicals from which they are formed. When the samples are dissolved in an acidic buffer, the hydroperoxides react with the transition metal (mainly iron) ions released from the proteins in the acidic medium and are converted to alkoxy and peroxy radicals. These newly-formed radicals oxidize an additive aromatic amine (N, N-diethyl-para-phenylen-diamine) and cause the formation of a relatively stable colored cation radical that is spectrophotometrically detectable at 505 nm. The results are expressed in arbitrary units (U. Carr) with one unit corresponding to 0.8 mg/L of hydrogen peroxide. According to manufacturer’s indications and previous preclinical and clinical studies (9, 10), ROS levels >32 mg H2O2/L are considered indicative of high oxidative stress; values between 27 and 32 mg H2O2/L correspond to a moderate oxidative stress, while ROS serum levels <27 mg H2O2/L indicate a low oxidative stress.

Data are expressed as the mean ±1 standard deviation and analyzed using the Prism statistical package (GraphpadInstat, version 3). Continuous variables were evaluated by using the one-way analysis of variance followed by paired t tests, while categorical data were calculated using the Fisher’s exact probability test or χ2-test, when appropriate. P values <0.05 were considered statistically significant.

■ RESULTS

At baseline all patients had a high disease activity (mean DAS28-CRP = 6.4±0.9) and during the treatment we observed a significant improvement in DAS28 both after 24 weeks (3.5±1.2) and 52 weeks (3.3±1.1) (p<0.001). After 24 weeks of treatment, 22 patients reached a good response, 14 patients had a moderate re-

![Figure 1 - Association of oxidative stress in rheumatoid arthritis patients with disease activity (A) and disability (B). Oxidative stress evaluated by reactive oxygen species serum levels: high >32 mg H2O2/L; moderate 27-32 mg H2O2/L; low <27 mg H2O2/L. Disease activity according to disease activity score on 28 joints/C-reactive protein (DAS28-CRP): high >5.1, moderate 3.2-5.1; low <3.2. Disability assessed by health assessment questionnaire disability index (HAQ-DI): high >1.0, moderate 0.5-1.0; low <0.5. A) χ2=247; p<0.001; B) χ2=138; p<0.001.](image)
response and 4 patients had no response according to the EULAR definitions. The last 4 patients dropped out from the study and were switched to other treatments. After 52 weeks, 5 more patients reported a good EULAR response. The HAQ significantly decreased from 1.8±0.7 at baseline to 0.9±0.4 and 0.7±0.5 after 24 and 52 weeks of treatment, respectively (p<0.001).

As to the ROS serum levels of RA patients, at baseline we observed a mean ROS of 33.2±10 mg H2O2/L, with a high oxidative stress in 23 patients (>32 mg H2O2/L), and a moderate oxidative stress in 17 patients (ROS <32 and >27 mg H2O2/L).

After 24 weeks of anti-TNFα treatment, the mean ROS serum levels dropped to 29.5±7 mg H2O2/L (p=0.02 vs baseline). The 20 patients who reached a low disease activity (DAS28-CRP <3.2) had a low oxidative stress status (ROS <27 mg H2O2/L), while the 4 patients who had no EULAR response had a high oxidative stress (ROS >32 mg H2O2/L).

After 52 weeks of treatment, ROS serum levels were 29.3±9 mg H2O2/L (p=0.02 vs basal), 26 out of 36 patients (72%) had a low disease activity (DAS28-CRP <3.2) and a low oxidative stress status (ROS <27 mg H2O2/L).

Lastly, we also observed a significant association between the oxidative stress evaluated by measuring the circulating ROS serum levels and the disease activity according to DAS28-CRP (χ²=247; p<0.001) and the functional status of RA patients assessed by the HAQ-DI (χ²=138; p<0.001) (Fig. 1).

**DISCUSSION AND CONCLUSIONS**

Our study confirmed the correlation between circulating ROS levels and disease activity in RA patients. However, the mechanisms responsible for the onset of RA remain unclear. Smoking is considered to be one of the most important extrinsic risk factors with a role in its development and severity (11). Recent studies demonstrated the interrelation between smoke, oxidative stress, inflammation, autoantibody formation and epigenetic changes in RA (12). ROS play a significant role in the progressive joint destruction both upstream and downstream to the nuclear factor kappa B (NF-κB) and TNF-α pathways, which are located at the center of the inflammatory response (13). Moreover, the increased oxidative stress is actually considered the main key driver in RA extrajoint morbidity, mainly in accelerated atherosclerosis, which may led to increased cardiovascular disease and mortality. The immune response, via cytokines and chemokines that attract monocytes, plays a role in these alterations ranging from the formation and stabilization of the atherosclerotic plaque to its progression and rupture (14). The control of the systemic inflammation is a therapeutic goal which not only translates in the remission of musculoskeletal symptoms, but also in an improvement of the patient overall health. Recent biologic immunosuppressive therapies targeting pro-inflammatory cytokines proved to be able to control disease activity and halt the progressive joint destruction (15). TNF-α inhibitors exert an anti-oxidative stress activity interfering with the pleiotropic effect of this cytokine. TNF-α stimulates NAD(P)H oxidases activity and increases intracellular hydrogen peroxide. The up-regulation of TNF-α impairs NO bioavailability and contributes to the development of mitochondrial oxidative stress (16). Lastly, the activation of the Caspase cascade induced by TNF-α demonstrated to promote endothelial apoptotic cell death which, along with increased oxidative stress and vascular inflammation, increases the cardiovascular risk (17). The beneficial action of anti-TNF-α drugs on endothelial cells seems to be ascribable to their excellent anti-inflammatory properties (18), yet their metabolic and cardiovascular effects are still unclear. In the early 2000s, some trials with anti-TNF-α raised some concern about its cardiovascular safety, worsening previous heart failure, while the anti-interleukin 6 treatment demonstrated to modify the lipid pattern by creating an atherogenic profile. On the other hand, data from National registries of RA patients seem to
demonstrate a reduction in cardiovascular events in responders to the biological treatment (19). High levels of ROS are associated with obesity, cardiovascular diseases and atherosclerosis. Only few studies investigated the effects of TNF-α blocking therapy on the oxidative stress. In accordance with Biniecka et al. (20) and Kageyama et al. (21), we observed that TNFα antagonism is able to reduce oxidative stress in responders. A lack of effective control of the disease activity with CCS and DMARDs results in a state of high inflammation and concurrent high circulating levels of ROS. On the contrary, the addition of anti-TNF-α agents led to a good disease control and consequent reduction of ROS levels and oxidative stress.

Our demonstration of a reduction in circulating ROS levels during an anti-TNF-α treatment, subcutaneously administered in combination with CCS and/or DMARDs, may explain the ability of these agents to reduce cardiovascular morbidity and mortality in RA patients who achieve a good disease control.

Previous studies highlighted the potential role of increased lipid peroxidation and decreased enzymic and non-enzymic antioxidants in RA, suggesting a very important role of oxidative stress in the pathogenesis of the disease. A previous study by Jacobson et al. (22) failed to demonstrate a correlation between plasma glutathione peroxidase concentrations and the severity of the disease measured using the HAQ-DI. To our knowledge, this is the first study that observed a significant correlation between the oxidative stress evaluated by the d-ROM-test and the severity of the HAQ-DI.

RA patients with more severe symptoms of pain and joint inflammation at disease onset have an increased risk of falling in the most disabled group of patients later, with a direct relationship between joint and systemic inflammation and loss of functional ability (23). The early management of systemic and joint inflammation with an effective treatment, as observed in good responders, and above all the resulting improvement in tender and swollen joints count of the composite indices of disease activity lead to a good control of pain and function with better outcomes. Therefore, the improvement of oxidative stress observed in the patients achieving the DAS28-CRP low disease activity might also have an impact on the degree of disability with an improvement in HAQ-DI.

In conclusion, the inhibition of TNF-α is able to control the disease activity and reduce the oxidative stress and has a positive impact on the disability of RA patients. This finding may explain the systemic effects of anti-TNF-α agents and justify the initiation of an early treatment to prevent the extra-articular manifestations of the disease (i.e., cardiovascular comorbidity). As a whole, in patients with RA, the correlation between DAS28-CRP and ROS levels suggests that the measurement of oxidative stress may serve as a biomarker to monitor disease severity in RA.

Key messages
- Rheumatoid arthritis patients with active disease have high oxidative stress.
- Good disease control with TNF-α-inhibitors can reduce oxidative stress.
- Oxidative stress correlates with DAS28-CRP and HAQ-DI.

Conflict of interest: the authors declare no conflict of interest.

REFERENCES


