The pitfalls of using the 2010 American college of rheumatology/European league against rheumatism (acr/eular) classification criteria for diagnosis of rheumatoid arthritis

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Abstract
We report a case of a 63-year-old man with a history of monoarthritis, thought to have Rheumatoid Arthritis (RA) and treated with disease-modifying antirheumatic drugs (DMARDs). No clinical or serological improvement was observed. After five months synovectomy was performed. A foreign body was recovered, and synovial fluid analysis revealed a rare fungal infection, Phaeoacremonium sp. This is an educational case that reveals potential pitfalls of the specificity of using the current classification criteria for Rheumatoid Arthritis (RA) as a diagnostic tool in patients presenting with monoarthritis and highlights the need to continually reassess the diagnosis when there’s inadequate response to treatment.

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Introduction
In the absence of a gold standard test, the diagnosis of Rheumatoid Arthritis (RA) can be challenging. The first classification criteria from the American College of Rheumatology came in 1987, and it had low sensitivity and specificity for diagnosing early-stage disease [1,2]. Twenty-three years later, the ACR collaborated with the European League Against Rheumatism (EULAR) to come up with new criteria that are more sensitive and specific in diagnosis than the 1987 criteria [3-5].

Case presentation
A 63-year-old Caucasian male was referred to the rheumatology clinic for a four-year history of left fourth finger pain. The pain was in the proximal interphalangeal (PIP) joint, he noted worsening in the past 10 months, it was non-radiating, asymmetrical and worst with movement. It was accompanied by morning stiffness. The patient rated the pain as 8 out of 10 in severity. There was no numbness or tingling.

Past medical history included dyslipidemia and left hip osteoarthritis. Family history was irrelevant except that his father had Psoriasis. The patient doesn’t smoke or drink alcohol. He does gardening. He was taking Lovastatin, ASA, and Ibuprofen-Famotidine at that time. Review of system was noncontributory. On physical exam, the proximal interphalangeal (PIP) joint of the left fourth finger showed tenderness and limited range of motion without erythema. Peripheral pulses and capillary refill were brisk and normal. Sensation was intact. There was no muscles atrophy.

X-ray (Figure 1) showed periarticular erosions along the medial and lateral aspect of the head of the PIP joint, with severe fusiform soft tissue swelling at the level of PIP. This indicated inflammatory changes in the joint. Laboratory testing showed elevated CRP 4.53 mg/L (0-3 mg/L) and elevated anticitrullinated protein antibodies (ACPA) 225 EU/ml (0-20 EU/ml). Rheumatoid Factor (RF) was within normal range, 11.2 IU/ml (0-15 IU/ml). Other serological markers and laboratory investigations were within normal limits.

The Patient was diagnosed with Rheumatoid Arthritis. He was started on Methotrexate (MTX) 15mg/week and received a Triamcinolone 10 mg intraarticular injection in the joint. The patient had significant improvement with the local injection for five weeks, but he had recurrence of the pain and swelling after eight weeks. A decision was taken to start the patient on Etanercept 50mg/ml/week and increase the MTX dose to 20 mg/week. There was no significant improvement at the six-week follow up despite normalization of CRP to 2.4 mg/L. ACPA was rechecked and was at a higher titer of >250 EU/ml. An MRI (Figure 2) was ordered to rule out alternative diagnosis, it showed large joint effusion involving the ring finger PIP joint with synovial enhancement associated with marrow edema and enhancement. There was surrounding soft tissue swelling as well. The patient was referred for a synovectomy which was performed a month later.

A foreign body was discovered during the surgery. The patient had a 0.4 x 0.3 x 0.2 cm3 splinter. Histopathology showed focal acute inflammation, fibrin deposition on the surface as well as intense lymphoplasmacytic infiltrates and focal foreign body granulomatous...
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The 2010 American College of Rheumatology/European League Against Rheumatism classification criteria for RA (score-based algorithm: add score of categories A–D; a score of 6/10 is needed for classification of a patient as having definite RA).

<table>
<thead>
<tr>
<th>Score</th>
<th>A. Joint involvement</th>
<th>B. Serology (at least 1 test result is needed for classification)</th>
<th>C. Acute-phase reactants (at least one test result is needed for classification)</th>
<th>D. Duration of symptoms</th>
</tr>
</thead>
<tbody>
<tr>
<td>6/10</td>
<td>1 Large Joint</td>
<td>Negative RF and negative ACPA</td>
<td>Normal CRP and normal ESR</td>
<td>&lt;6 weeks</td>
</tr>
<tr>
<td>5</td>
<td>≥1 Joint</td>
<td>Low-positive RF or low-positive ACPA</td>
<td>Abnormal CRP or abnormal ESR</td>
<td>≥6 weeks</td>
</tr>
<tr>
<td>4</td>
<td>≥2 Large joints</td>
<td>High-positive RF or high-positive ACPA</td>
<td></td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>≥1-3 small joints</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>2</td>
<td>≥4–10 small joints (with or without involvement of large joints)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>1</td>
<td>≥10 joints (at least one small joint)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>0</td>
<td>0</td>
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Figure 1. X-Ray image showing severe inflammatory changes and erosions

Figure 2. MRI showing large joint effusion involving the ring finger PIP joint with synovial enhancement associated with marrow edema and enhancement. Soft tissue swelling is noted as well

Discussion

Rheumatoid Arthritis (RA) is a potentially disabling, systemic autoimmune inflammatory disease of unknown etiology. It affects about 1% of the population and if not treated or if refractory to therapy it may result in joint destruction, disability, and loss of daily function and employment. Recent data suggest that early diagnosis and initiation of disease-modifying antirheumatic drugs (DMARDs) may reduce morbidity and mortality in RA patients [6]. However, the diagnosis of early RA has been a challenge for decades and clinical trials are still restricted by the difficulty of diagnosing early cases of RA because of lack of more accurate diagnostic criteria [3]. Therefore, the establishment of reliable and accurate diagnostic tools for early disease is of a paramount importance. Specially, when intensive treatment early in the disease may change or modify the natural history of RA and prevent destructive bony erosions [7].

The 1987 ACR classification criteria were used to diagnose RA for many years. Even though they had pooled sensitivity and specificity of (77-80% and 33-77%) respectively in early disease and (79-80% and 90-93% respectively) in established disease, they were not meant to be used for diagnosis [1]. In 2010 a joint working group of the ACR and EULAR was therefore formed to develop a new approach for classification of RA. The working group investigated the factors that were thought to discriminate people with undifferentiated arthritis based on the likelihood of developing persistent and/or erosive disease which is the new paradigm of RA.

The ACR/EULAR report states that for RA to be classified as definite it requires the confirmed presence of synovitis in at least one joint, the absence of an alternative diagnosis for the observed arthritis, and a total score of at least 6 to be deemed RA [3-5] (Table 1). Our patient scored 7. It is true that the 2010 criteria were initially formulated for classification and research purposes to recruit patients for clinical trials and testing of early RA treatment and it is true that the presence of ACPA antibodies is 94.5% specific for the diagnosis of RA [8]. Based on that evidence, the patient was diagnosed with RA and the decision was made to initiate treatment.

Of interest, psoriasis which can cause monoarthritis and elevated ACPA [9], was considered as a diagnosis, specially that the patient had a positive family history of psoriasis. Nevertheless, the treatment with MTX and Etanercept should have alleviated the patient’s symptoms if that was the case.

Unfortunately, an unknown wooden splinter had introduced an opportunistic fungus, Phaeoacremonium sp, that’s found in soil and woody plants [10], and mainly infects the immunocompromised. This infection can be serious and fatal if disseminated [11]. The patient’s symptoms improved after a local steroid injection, however, it is not uncommon for septic arthritis to improve after a local steroid injection [12].

More studies outlining the diagnosis of early RA should be done. Fitting the ACR/EULAR 2010 criteria shouldn’t be necessary.
be conclusive for a diagnosis of early RA or RA with a single joint involvement. In monoarthritis, it is recommended that the joint should be aspirated, and infections must be ruled out, regardless of the duration of symptoms and/or if they fit into any other diagnoses. Otherwise, drastic complications may occur. Also, if appropriate it has been suggested that High-Resolution Ultrasoundography, can detect plant thorns [13] and foreign bodies and maybe they should be used as a first study to do in monoarthritis because of their practicality and convenience.

**Limitations**

The patient failed to mention any history of trauma, which he had. Had he mentioned it, a better differential diagnosis could have been made and perhaps synovial fluid could have been aspirated and analyzed.

**Conclusion**

This case points out a potential pitfall in the 2010 ACR/EULAR classification criteria for RA, when used as a diagnostic tool for diagnosing RA in patients with monoarthritis.

Although some patients can present a challenge to treat, this case reinforce the need to continually reevaluate patients when they do not respond to what would be an appropriate treatment.

**Conflict of Interest**

None Declared

**Disclosures**

None Declared

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