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Guest Editorial

The two extremes of rheumatic diseases possible appropriate definition and their bearing

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It is a famous saying that “extremes” of anything are precarious. When it comes to Rheumatic Diseases, the two extremes of the disease can be identified –

1. “Very very early stage”, preferably called “pre-rheumatic stage”
2. “Refractory to treatment”. Practically denotes refractory to various treatment protocols.

Identification of the first extreme, “the pre-clinical stage”, helps in deciphering the future likelihood of the disease. At this stage, therapeutic intervention is likely to either abort the disease or achieve an early remission.

Identification of the other extreme, when the disease can be defined as refractory to treatment or “difficult to treat” (D to T) rheumatoid arthritis.

These symptomatic patients of rheumatoid arthritis reflect a complex interplay of disease. This stage is a stimulus for opening up new vistas of therapy including “alternative medicine”.

The principal goal in the management of rheumatoid arthritis is reaching or achieving sustained remission or produce minimum or low disease activity. The earliest phases of rheumatoid arthritis have been designated variously as probable rheumatoid arthritis, very early rheumatoid arthritis, pre-clinical rheumatoid arthritis,

autoantibody-positive arthralgia, and early undifferentiated arthritis progressing to rheumatoid arthritis. To avoid confusion, the term “pre-rheumatoid arthritis” has been coined for the evolving stage of the disease, before the clinical appearance of rheumatoid arthritis. The presence of abnormalities in the immune function and immune responses characterize this stage.

The environmental factors’ interplay with the genetic factors sequentially. This is suggested to initiate and propagate the required autoimmune process to generate the disease, rheumatoid arthritis. Finally, the inflammatory response of the tissues starts producing symptoms.

The disease-related autoantibodies like anticitrullinated peptide antibodies (ACPA) develops and sometimes also rheumatoid factors (RF), much before clinical features of tissue inflammation appear. Minimal features developing at the earliest labels them as unclassified or non-specific before the symptoms, signs fully bloom into rheumatoid arthritis.

These ‘at risk’ individuals pass through different stages before the disease appears with its classical picture.

Stage I – Interplay of environmental and genetic risk factors.

Stage II – Production of autoantibodies like ACPA and sometimes RF.

Stage III - Arthralgia / joint stiffness but no clinical arthritis.

Stage IV - Arthritis in one / two joints – early

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/ undifferentiated arthritis intermittent appearance is pronounced foalindronic rheumatism.

Stage V – Some patients develop into typical classical established rheumatoid arthritis.

The “pre-rheumatoid arthritis” can be labeled before any stage of the disease but only retrospectively in a patient, who has subsequently developed rheumatoid arthritis.

1. The environmental risk factors of rheumatoid arthritis need to be discussed first because these contributing factors are preventable like:
 1. Infection
 2. Periodontitis
 3. Smoking
 4. Dietary factors
 5. Obesity
 6. Hormonal factors
2. Genetic risk factors - HLADR4 is the genetic link in about 70% of rheumatoid arthritis patients. This susceptibility associated with the DRβ1 allele – the third hypervariable region – is called SE (susceptibility epitope). PTPN 22 and PADI-4 (Protein tyrosine phosphatase 22 and peptidyl arginase deiminase) are the other two genes that carry the two-fold risk of developing rheumatoid arthritis, apart from other genes carrying lower risk.
3. Symptoms without clinical evidence of arthritis - Arthralgia or joint stiffness with positive rheumatoid factor and/or ACPA can unusually be seen without any evidence of arthritis. After the passage of sometimes, they progress to classical rheumatoid arthritis, denoting it to be a ‘pre-RA phase’.
4. Systemic Autoimmunity Associated with RA – Asymptomatic patients having positive rheumatoid factor and/or ACPA are grouped under this heading. This is based on the studies that antibodies related to RA are present in the system before the diagnosis of RA, making it abundantly clear that the presence of rheumatoid factor and ACPA is a highly specific predictor of the oncoming RA.
5. Undifferentiated Arthritis - There is the involvement of more than monoarticular affection without fulfilling the criteria for RA or any other connective tissue disease, finally 30-40% progress to classical RA in long-term follow-up. This “diagnosis of exclusion” does not satisfy the classification criteria of a well-recognized rheumatic disease, still has the potential for the development of persistent inflammatory arthritis.

Screening individuals with a high risk of developing RA in the future should selectively be used amongst first-degree relatives of RA patients, autoantibody-positive individuals, twins of RA patients. This screening can also cover the population with high disease prevalence. Though these studies prove costly, they can evolve into a

management program where the disease can be arrested or aborted even before it becomes clinically evident. Such screening programs can evolve into preventive strategies. Symptomatic patients require to be managed effectively with early achievable remission or low disease activity.

1. The Other Extreme

A significant proportion of patients with rheumatoid arthritis continue to remain symptomatic even after the management of appropriate recommendations. They have been coined “difficult-to-treat” (D2T) RA. EULAR steering committee resolved this terminology based on a survey amongst international rheumatologists.

The undermentioned three criteria have been agreed to define the mandatory components within the framework of “difficult-to-treat” RA:

1. Failure of $\geq$ biological DMARDs/ tsDMARDs even after treatment as per EULAR recommendations after failure of csDMARDs (unless contraindicated).
2. Presence of at least one –
 - a. Moderate disease activity at least.
 - b. Presence of active disease - Signs and/or symptoms.
 - c. Inability to taper glucocorticoid treatment.
 - d. Presence of rapid radiographic progression.
 - e. Compromised quality of life due to RA symptoms
3. Problematic management of signs and/or symptoms as perceived by rheumatologists and/or patients.

A clear definition has been lacking for these patients eluding remission or low disease activity despite the addition of newer drugs and new management approaches. Different terms have been in use for these patients like severe, refractory, resistant to multiple drugs or treatments, complex, and established. Difficult-to-treat is a widely accepted term in pulmonology, psychiatry, and cardiology, hence, was accepted by the EULAR task force.

The clinical challenge of managing difficult-to-treat RA will require practical management recommendations. Several factors including persistent inflammatory activities due to the resistance of the disease to DMARDs, limited drug options due to adverse drug reactions, and/or co-morbidities precluding the use of DMARDs or non-adherence to treatment, may complicate the management of these patients. Concomitant diseases like psychosocial factors, fibromyalgia, and osteoarthritis can result in non-inflammatory pain mimicking inflammatory activity. EULAR management recommendations are still under development but hopefully are likely to cover both underlying inflammatory and non-inflammatory factors. The definition encompassed:

1. Treatment failure history
2. Characterization of active / symptomatic disease

3. Clinical perception

Acceptable glucocorticoid dose for chronic use remains a matter of discussion but there is no denying the fact that a significant number of RA patients are treated with long-term glucocorticoids. Current EULAR recommendations consider using glucocorticoid on initiating or changing CsDMARDs with simultaneous emphasis on tapering glucocorticoid as rapidly as clinically feasible. The risk of harm is generally low at long-term doses of ≤ 5 mg prednisone equivalent/day as perceived by the task force in charge of evaluating the risk of long-term GC therapy. Inability to taper glucocorticoid treatment below 7.5 mg/day prednisone or equivalent has been listed as criteria by this EULAR task force with its limitations.

Achieving sustained remission or low disease activity following current EULAR recommendations is the principal goal of RA management. A newer management protocol should necessarily be developed for these difficult-to-treat RA patients. Robust and consistent identification of these difficult-to-treat patients has been achieved by this EULAR definition requiring further work to provide detailed recommendations for management.

2. Further Reading

1. Raza k, klareskog L, Holers VM .Predicting and preventing the development of rheumatoid arthritis Rheumatology(Oxford) 2016;55:1-3.
2. Gerlag DM, Raza K, van Baarsen LG, Brouwer E, Buckley CD, Burmester GR, et al. EULAR recommendations for terminology and research in individuals at risk of rheumatoid arthritis: report from the Study Group for Risk Factors for Rheumatoid Arthritis. Ann Rheum Dis.2012; 71:638-41.
3. Turk SA, van Beers-Tas MH, van Schaardenburg D. Prediction of future rheumatoid arthritis. Rheum Dis Clin North Am 2014; 40:753-70.
4. Van Steenberg HW, Huizinga TW, van der Helm-van Mil AH. The preclinical phase of rheumatoid arthritis: What is acknowledged and what needs to be assessed? Arthritis Rheum 2013; 65:2219-32.
5. Sugiyama D, Nishimura K, Tamaki K, Tsuji G, Nakazawa T, Morinobu A, et al. Impact of smoking

as a risk factor for developing RA: A meta-analysis of observational studies. Ann Rheum Dis 2010; 69 70-81.

6. Karlson EW, Deane K. Environmental and gene-environment interactions and risk of rheumatoid arthritis. Rheum Dis Clin North Am 2012;38:405-26
7. Binoy JP, Geetha P, Shanu PM. Clinical profile and long-term sequel of chikungunya fever. Indian J Rheumatol 2011; 6(1):12-19.
8. Joseph R, Rajappan S, Nath SG, Paul BJ. Association between chronic periodontitis and rheumatoid arthritis: a hospital-based case-control study. Rheumatology (Oxford) 2014; 53:526-31.
9. De Hair MJ, Landewe RB, van de Sande MG, van Scharrendburg D, van Basarsen LG, Gerlag DM, et al. smoking and overweight determine the likelihood of developing rheumatoid arthritis. Ann Rheum Dis 2013; 72:1654-8.
10. van der Helm-van Mil AH, le Cassie S, van Dongen H, Breedveld FC, Toes RE, Huizinga TW. A prediction rule for disease outcome in patients with recent-onset undifferentiated arthritis. how to guide individual treatment decisions. Arthritis Rheum 2007; 56:433-40.
11. Lahri M, Morgan C, Symmons DP, Bruce IN. Modifiable risk factors for RA: prevention, better than cure? Rheumatology (Oxford).2012; 51:499-512.

3. Conflict of Interest

The authors declare that there are no conflicts of interest in this paper.

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