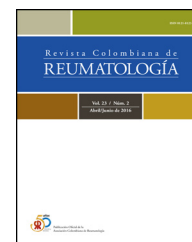




Revista Colombiana de REUMATOLOGÍA

www.elsevier.es/rcreuma



Original Investigation

Frequency of anergy in a group of patients with rheumatoid arthritis on immunosuppressive therapy



S.A. Vallejo*, H.S. Basallo, M. Narvaes, Y.F. Medina, G. Quintana-López

Department of Internal Medicine, National University of Colombia, Bogotá, Colombia

ARTICLE INFO

Article history:

Received 17 February 2020

Accepted 13 June 2020

Available online 25 August 2020

Keywords:

Anergy

Cutaneous anergy

Rheumatoid arthritis

Latent tuberculosis

ABSTRACT

Background: Tuberculin is the globally accepted delayed cutaneous hypersensitivity test for the diagnosis of latent tuberculosis. The alteration of cellular immunity induced by disease-modifying drugs used in rheumatoid arthritis may give a false negative result, also known as cutaneous anergy. There are no studies that determine the frequency of anergy in patients with rheumatoid arthritis and on immunosuppressive therapy.

Objective: To determine the frequency and possible factors associated with cutaneous anergy in a group of patients with rheumatoid arthritis and on immunosuppressive therapy.

Methods: Cross-sectional analytical observational study including 100 patients with rheumatoid arthritis on immunosuppressive therapy. They were tested for delayed cutaneous hypersensitivity with tuberculin, and a control test with tetanus toxoid. The non-reactivity of both tests was defined as anergy.

Results: The overall frequency of cutaneous anergy was 9% (n = 11). It occurred in 33% of men versus 6% of women. The mean age was 57 years, and 89% were over 50 years-old. Being female behaved as a protective variable for the generation of anergy, OR 0.795 [95% CI, 0.658 - 0.959, P<.05]. All patients with anergy were being treated with corticosteroids, 44% with methotrexate, and 33% with biological therapy. Treatment with moderate to high dose prednisone and biological therapy were independently associated as risk factors for presenting with anergy, OR 1.044 [95% CI, 1.008-1.080 P<.05] and OR 1.096 [95% CI, 1.016-1.182, P<.05], respectively. The overall positivity for tuberculin was 13%. Symptoms associated with disease activation were present in 38% of these. All cases (n= 1) of confirmed active tuberculosis were excluded.

Conclusions: The high prevalence of cutaneous anergy in patients with RA in the present study, and the evidence presented here, supports the recommendation of a second diagnostic test (tuberculin booster or Interferon-Gamma Release Assays) for the diagnosis of latent TB in patients with RA on immunosuppressive therapy.

© 2020 Asociación Colombiana de Reumatología. Published by Elsevier España, S.L.U. All rights reserved.

* Corresponding author.

E-mail address: savallejoa@unal.edu.co (S.A. Vallejo).

<https://doi.org/10.1016/j.rcreu.2020.06.004>

0121-8123/© 2020 Asociación Colombiana de Reumatología. Published by Elsevier España, S.L.U. All rights reserved.

Frecuencia de anergia en un grupo de pacientes con artritis reumatoide y terapia inmunosupresora

R E S U M E N

Palabras clave:

Anergia

Anergia cutánea

Artritis reumatoide

Tuberculosis latente

Antecedentes: La tuberculina es la prueba de hipersensibilidad cutánea tardía mundialmente aceptada para el diagnóstico de tuberculosis latente. La alteración de la inmunidad celular inducida por los fármacos modificadores de la enfermedad utilizados en la artritis reumatoide puede dar un resultado falso negativo, también conocido como anergia cutánea. No hay estudios que determinen la frecuencia de anergia en pacientes con artritis reumatoide y terapia inmunosupresora.

Objetivo: Determinar la frecuencia y los posibles factores asociados con la anergia cutánea en un grupo de pacientes con artritis reumatoide y terapia inmunosupresora.

Métodos: Estudio observacional analítico transversal que incluyó a 100 pacientes con artritis reumatoide con terapia inmunosupresora. Se les realizó una prueba de hipersensibilidad cutánea tardía con tuberculina y una prueba de control con toxoide tetánico. La no reactividad de ambas pruebas se definió como anergia.

Resultados: La frecuencia general de anergia cutánea fue del 9% ($n = 11$). Ocurrió en el 33% de los hombres versus el 6% de las mujeres, la edad promedio fue de 57 años y el 89% tenía más de 50 años. El sexo femenino se comportó como una variable protectora para la generación de anergia (OR 0,795; IC 95%: 0,658-0,959; $p < 0,05$). Todos los pacientes con anergia usaron corticosteroides, el 44% fue tratado con metotrexato y el 33% con terapia biológica. El tratamiento con dosis de moderadas a altas de prednisona y terapia biológica se asoció de manera independiente como factor de riesgo para la presentación de anergia: OR 1,044 (IC 95%: 1,008-1,080; $p < 0,05$) y OR 1,096 (IC 95%: 1,016-1,182; $p < 0,05$), respectivamente. La positividad general para la tuberculina fue del 13%. Los síntomas asociados con la activación de la enfermedad estaban presentes en el 38% de ellos. Se excluyeron todos los casos de tuberculosis activa confirmada ($n = 1$).

Conclusiones: La alta prevalencia de anergia cutánea en pacientes con artritis reumatoide en el presente estudio y la evidencia presentada respaldan la recomendación de una segunda prueba de diagnóstico (refuerzo de tuberculina o IGRA) para el diagnóstico de tuberculosis latente en pacientes con artritis reumatoide y terapia inmunosupresora.

© 2020 Asociación Colombiana de Reumatología. Publicado por Elsevier España, S.L.U.

Todos los derechos reservados.

Introduction

In rheumatoid arthritis (RA), synthetic and biological immunosuppressive therapy are the cornerstone of treatment. This type of therapy might carry a greater risk of reactivation of latent chronic infections such as tuberculosis (TB) up to four times above the population without these treatments.^{1,2} The majority of people control the infectious process after exposure, infection and subsequent dissemination with persistent latent viable germs contained in granulomas without presenting clinical manifestations.^{3,4}

According to the latest global report, Colombia is a country with an intermediate incidence of 25 to 49 TB cases per 100,000 inhabitants and it is estimated that one third of the population may have latent TB.^{5,6} Between 5 to 15% of people with latent TB develop active infection during life.⁷ This depends on the virulence of the bacteria and immunity of the host.^{4,8,9} A relationship has been found with the use of synthetic, biological and corticosteroid immunosuppressive therapy with the transformation of a latent phase to an active phase of TB.^{4,8,9} Specifically in RA, there is evidence of TB reactivation with

disease-modifying medications, in particular tumor necrosis factor alpha (iTNF- α) inhibitors.¹⁰⁻¹²

The World Health Organization (WHO) and the Center for Disease Control (CDC) recommend the detection of latent disease in high-risk patients with the intention of treating any positive test.^{4,13} Tests should be practiced systematically in people infected with HIV, in contact with those infected with pulmonary TB, prior to initiating biological therapies, people on dialysis, solid organ recipients or hematological transplantation, among others.¹³⁻¹⁹ Indirect tests are used for diagnosis by measuring the immune response in vivo such as tuberculin, or in vitro, with the interferon gamma release assay (IGRAs). There is no quality evidence that one is superior to the other.^{20,21}

The tuberculin skin test is based on the principle that *Mycobacterium Tuberculosis* infection produces a delayed hypersensitivity reaction mediated by T lymphocytes in response to certain antigens present in extracts from filtered cultures called tuberculins. These purified derived proteins (PPD) produce skin induration after vasodilation and chemotaxis of inflammatory cells.^{4,22-24} The standard dose is 0.1 ml intradermally (concentration of 5 tuberculin units) per

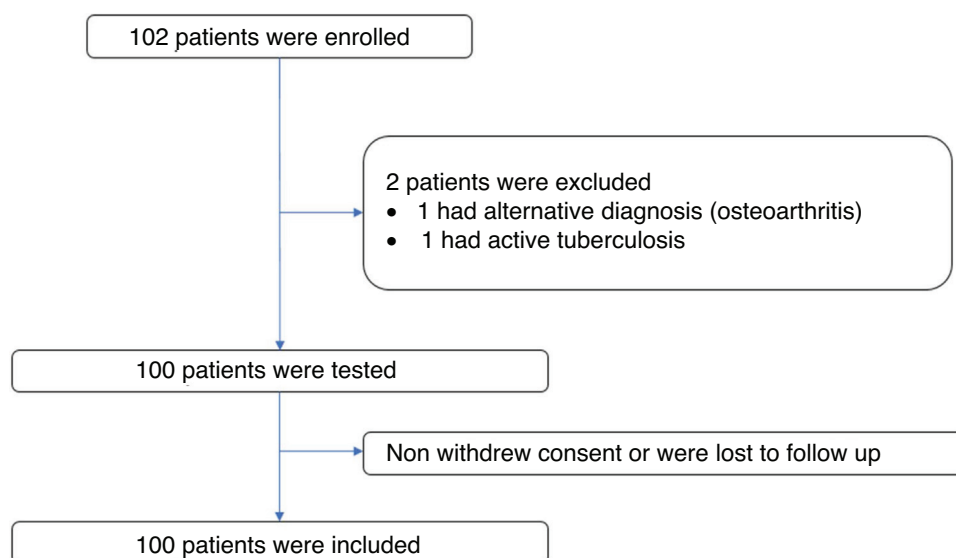


Figure 1 – Enrollment and follow-up.

single puncture. It is done in the forearm area, which should be free of lesions and preferably away from venous vascular areas.^{23,25,26} The reading should be carried out between 48 and 72 hours after the puncture for visual evaluation and / or palpation. The use of the Sokal method is recommended to mark the indurated area.^{23,25,26} It is a safe test that rarely generates allergic reactions.

The sensitivity of the test is variable and the frequency of false negatives can reach 50% according to some reports in patients with different types of immunosuppression.²⁷⁻³² On the other hand, there are conditions that may occur with induration in the absence of tuberculous infection such as non-tuberculous mycobacterial exposure.^{33,34}

The tuberculin test depends on the integrity of the cellular immune system that can be altered by noxas such as the use of immunosuppressive therapy. The absence of reaction in a previously exposed patient is known as anergy and is objectified by the inability to express delayed hypersensitivity to skin tests of common antigens.³⁵⁻³⁸

Anergy has been characterized in people with HIV infection, dialysis, diabetes, malnourished population, among others.^{22,27-32,39-41} It is also found in healthy people where IL-10 producing T cells are constitutively activated.⁴⁰⁻⁴³

It is plausible that pharmacological immunosuppression in RA causes anergy by altering cellular immunity. This has encouraged different scientific societies to recommend conducting a tuberculin booster two weeks later to improve sensitivity or another sequential alternative test (IGRAs).^{4,13-16,19,44} Thus, boosting the hypersensitivity response and reducing false negatives.^{14,19} Several studies have assessed these tests in groups of patients on dialysis, increasing the number of positive tuberculin by 12%^{17,45} and also in RA where the reinforcement increased positivity by 15%.^{17,18}

However, this recommendation has no scientific support as there are no studies that establish the prevalence of anergy in patients with RA with synthetic or biological immunosuppressive therapy. In our environment with difficulties in accessing

health services, the requirement of an additional test may delay the start of therapy for months.

In order to objectify anergy, ubiquitous and frequent antigens in the general population to which a healthy person should always react are used as skin tests. The negativity of these could be interpreted as anergy. The most frequently used antigens are those derived from candida, trichophyton, mumps virus, tetanus toxoid and streptokinase-streptodornase.^{36,38,46-57} The use of antigens such as tetanus toxoid and candidines have proven to be the diagnostic standard, other candidates historically used such as mumps virus, can lead to confusion and have been relegated from this type of study.⁵⁸ The interpretation of these tests is also due to the skin induration area such as tuberculin.⁵⁹⁻⁶¹

The frequency of late cutaneous hypersensitivity to tetanus toxoid in healthy subjects is 79% to 90% and has a good correlation with the leukocyte migration inhibition test, which makes it a valuable antigen in the evaluation of late hypersensitivity.⁶² The cutaneous reactivity to tetanus toxoid is independent of antibody titers for tetanus and no relationship has been demonstrated between immune response measures and the interval since the last booster immunization.^{62,63} Tetanus toxoid has proven useful in the evaluation of cellular immunity and as a marker of cutaneous anergy.^{40,42} It is a sensitive marker, has a low incidence of side effects and produces a slight but beneficial reinforcement of the serum antibody against tetanus toxoid.^{40,42,43} Additionally, vaccination does not elicit positivity in non-responders, which implies that the non-response is secondary to host factors rather than the absence of antigenic stimulation.⁶⁴

Induration diameter has varied through numerous studies that have used values greater than 2 mm,^{28,29,32,65-67} 3 mm²⁷ and 5 mm^{39,65,67} to be considered positive. In the present study we use a 5 mm cut as the diagnostic standard.

In general terms, the concomitant failure of the response of at least one skin control in patients who have tuberculin applied to them is defined as anergy.^{27,32,66,67} Whether there is a single negative result⁶⁸ or a negative result to all controls

Table 1 – Distribution of the sociodemographic aspects of the participants (n = 100).

	Absolute frequency	Relative frequency
Gender		
Male	12	12%
Female	88	88%
Age (median)	57	
20 to 30 years	5	5%
31 to 40 years	8	8%
41 to 50 years	15	15%
51 to 60 years	38	38%
61 to 70 years	26	19%
71 to 80 years	7	4%
Over 80 years	1	1%
Years from diagnosis		
Less than 2 years	15	15%
2 to 5 years	25	25%
6 to 10 years	28	28%
Over 11 years	42	42%
Diabetes	9	9%
Kidney disease	3	3%
Tuberculosis	3	3%
Malnutrition	2	2%
HIV	0	0%
Malignancy	3	3%
Methotrexate**	74	74%
Leflunomide**	46	46%
Antimalarial**	12	12%
Sulfasalazine**	18	18%
Prednisone**	69	69%
Prednisone >5 mg or combined therapy**	62	62%
Prednisone ≤5 mg monotherapy**	7	7%
Biological therapy**	16	16%

* Data are presented in median (interquartile range).

** Methotrexate 7.5 mg / week or more and / or leflunomide 10 mg / day or more and / or sulfasalazine 500 mg / day or more and / or prednisolone (or its equivalent) in doses greater than 5 mg day and / or biological therapy, all for more than three months

(if there are several) as in the case of the CMI Multitest manufactured by Merieux Institute that uses 7 antigens that it's no longer available.^{69,70} Currently, tests of one or two controls are used, such as the one used in this study, in which the tetanus toxoid was chosen as the most suitable.^{40,42}

The objective of this article is to estimate the frequency of anergy and latent TB in a group of patients with RA and immunosuppressive therapy in Bogotá during 2019. As well as describe the epidemiological, clinical and pharmacological characteristics that are related to this condition.

Materials and methods

Multicenter cross-sectional analytical observational study in which 102 adult volunteers with RA diagnosis were included by 2010 ACR / EULAR criteria with current use of immunosuppressive treatment for more than three months defined as: Methotrexate in doses greater than or equal to 7.5 mg / week and / or leflunomide at a dose greater than or equal to 10 mg / day and / or sulfasalazine at a dose greater than

or equal to 500 mg / day and / or prednisone (or its equivalent) at a dose greater than or equal to 5 mg per day and / or biological therapy. These subjects were captured in the rheumatology outpatient clinic based at the National University Hospital of Colombia and the integrated north subnet of health services of Bogotá during 2019. The ages of the participants ranged between 23 and 85 years and are stratified in 7 groups (20-30, 31-40, 41-50, 51-60, 61-70, 71-80 and > 80 years). Data on their age, weight, comorbidities, time of diagnosis of RA, immunosuppressive therapy and general symptoms were collected. Treatments such as azathioprine, cyclosporine, cyclophosphamide or mycophenolate mofetil were excluded. Patients with contraindications for skin tests, patients with active malignancy or other rheumatic disease were excluded. Patients with positive tuberculin tests were followed in the outpatient setting with chest x rays to rule out active disease. Patients with active or extrapulmonary pulmonary TB or in contact with confirmed or suspected TB and pregnant patients were also excluded.

Sampling

Patients were asked to participate in the study during the consultation or through a telephone call. Informed consent was signed explaining the procedures, advantages and risks of the intervention. A questionnaire was applied and background checks were done. After asepsis, two intradermal reagents were applied. The tuberculin is performed in the forearm by Mantoux technique with 5 UT in 0.1 ml of the solution, in the same way 0.1 ml of tetanus toxoid is administered 5 cm from the first. Induration reading was done 48 to 72 hours after application by certified personnel. Anergy is defined if the result of the induration is negative for both tests with a cut-off point less than 5 mm.^{27,28,32,39,65-68}

Latent TB is defined as a positive reaction in the tuberculin skin test in the absence of symptoms suggestive of active pulmonary infection (Cough or persistent dyspnea greater than 2 weeks, weight loss over 10% in the last 6 months, recurrent night sweating in the last 6 months or febrile peaks over 38 degrees in the last 6 months without identified infectious focus).⁷¹

Statistical analysis

For the descriptive analysis of the sociodemographic aspects and the findings of the skin tests of the patients, absolute distributions, relative distributions and summary indicators such as quartiles, interquartile range, maximum values and minimum values were used.

A non-probabilistic sampling was performed for the selection of the subjects to be included in the study (n = 102) to determine statistical estimates of association, with a power of 80%, a chance risk of 5% and a OR of 1.75 at minus based on an estimated anergy prevalence of 20%.⁷² The study groups were evaluated as nominal, continuous and percentile variables.

To determine the association of variables with the generation of anergy, Fisher's exact test or Chi square were used for categorical variables, as applicable. A confidence interval of 95% was taken into account. A p less than or equal to 0.05 was considered significant. The risk factors associated with

Table 2 – Distribution of the Anergy findings (n = 9) an No Anergy (n=81).

	Anergy		No Anergy	
	Absolute frequency	Relative frequency	Absolute frequency	Relative frequency
Gender				
Male	4	33	8	67
Female	5	6	83	94
Age groups				
20 to 30 years	0	0	4	5
31 to 40 years	1	11	7	9
41 to 50 years	0	0	14	17
51 to 60 years	5	56	33	41
61 to 70 years	1	11	25	31
71 to 80 years	2	22	5	6
Over 80 years	0	0	1	1
Years from diagnosis				
Less than 2 years	0	0	5	6
2 to 5 years	2	22	23	28
6 to 10 years	3	33	25	31
Over 11 years	4	44	38	47
Diabetes	1	11	8	10
Kidney disease	0	0	3	4
Tuberculosis	0	0	3	4
Malnutrition	0	0	2	2
HIV	0	0	0	0
Malignancy	1	11	2	2
Methotrexate**	4	44	70	86
Leflunomide**	1	11	45	56
Antimalarial**	1	11	11	14
Sulfasalazine**	1	11	17	21
Prednisone**	9	100	60	74
Prednisone >5 mg or combined therapy**	8	89	54	67
Prednisone ≤5 mg monotherapy**	1	11	6	7
Biological therapy**	3	33	13	16

** Methotrexate 7.5 mg / week or more and / or leflunomide 10 mg / day or more and/ or sulfasalazine 500 mg/ day or more and/ or prednisolone (or its equivalent) in doses greater than 5 mg day and/ or biological therapy, all for more than three months

Table 3 – Dependent variable logistic regression “Cutaneous anergy”.

Category	Regression coefficients	Standard error	Lower limit (LL)	Upper limit (UL)	Critical value t	p Value	Odds Ratio (OR)	LL (OR)	UL(OR)
Intercept	0.1702	0.2735	-0.3658	0.7061	0.6223	0.535587			
Age	0.0010	0.0027	-0.0044	0.0063	0.3472	0.729436	1.001	0.996	1.006
Gender	-0.2300	0.0962	-0.4186	-0.0415	-2.3912	0.019266	0.795	0.658	0.959
Weight	0.0012	0.0028	-0.0043	0.0068	0.4364	0.663808	1.001	0.996	1.007
Years from diagnosis	0.0005	0.0027	-0.0047	0.0058	0.2037	0.839127	1.001	0.995	1.006
Diabetes	0.0444	0.1120	-0.1750	0.2638	0.3964	0.692901	1.045	0.839	1.302
Kidney disease	-0.1623	0.1624	-0.4807	0.1560	-0.9995	0.320733	0.850	0.618	1.169
Previous tuberculosis	-0.0633	0.2138	-0.4822	0.3557	-0.2960	0.768063	0.939	0.617	1.427
Malnutrition	-0.1980	0.2021	-0.5941	0.1981	-0.9796	0.330371	0.820	0.552	1.219
Malignancy	0.1523	0.1694	-0.1797	0.4844	0.8992	0.371384	1.165	0.836	1.623
Methotrexate dosis	-0.0054	0.0042	-0.0135	0.0027	-1.3005	0.197370	0.995	0.987	1.003
Methotrexate time	-0.0041	0.0099	-0.0235	0.0153	-0.4128	0.680928	0.996	0.977	1.015
Leflunomide dosis	-0.0013	0.0035	-0.0081	0.0056	-0.3594	0.720329	0.999	0.992	1.006
Leflunomide time	-0.0194	0.0140	-0.0469	0.0082	-1.3777	0.172332	0.981	0.954	1.008
Antimalarial dosis	0.0008	0.0006	-0.0005	0.0021	1.2624	0.210672	1.001	1.000	1.002
Antimalarial time	-0.0215	0.0293	-0.0789	0.0359	-0.7336	0.465449	0.979	0.924	1.037
Sulfasalazine dosis	-0.0001	0.0001	-0.0003	0.0001	-0.8457	0.400351	1.000	1.000	1.000
Sulfasalazine time	-0.0103	0.0310	-0.0709	0.0504	-0.3322	0.740657	0.990	0.932	1.052
Prednisone dosis	0.0115	0.0102	-0.0085	0.0316	1.1269	0.263350	1.012	0.992	1.032
Prednisone time	0.0090	0.0117	-0.0138	0.0319	0.7740	0.441327	1.009	0.986	1.032
Biological therapy	0.0914	0.0386	0.0158	0.1670	2.3689	0.020382	1.096	1.016	1.182
Biological therapy dosis	-0.0002	0.0003	-0.0009	0.0004	-0.6841	0.495988	1.000	0.999	1.000
Biological therapy time	-0.0263	0.0931	-0.2089	0.1562	-0.2829	0.778048	0.974	0.811	1.169

Table 4 – Logistic regression combined variables.

Category	Regression coefficients	Standard error	Lower limit (LL)	Upper limit (UL)	Critical value t	p Value	Odds Ratio (OR)	LL (OR)	UL(OR)
Intercept	0.3903	0.5484	-0.6845	1.4651	0.7117	0.478858			
Age	0.0012	0.0026	-0.0039	0.0063	0.4653	0.643053	1.001	0.996	1.006
Gender	-0.2111	0.0899	-0.3873	-0.0349	-2.3484	0.021528	0.810	0.679	0.966
Weight	0.0004	0.0028	-0.0051	0.0059	0.1515	0.880004	1.000	0.995	1.006
Years from diagnosis	0.0007	0.0028	-0.0047	0.0061	0.2459	0.806429	1.001	0.995	1.006
Diabetes	0.0768	0.1076	-0.1341	0.2878	0.7138	0.477615	1.080	0.874	1.333
Kidney disease	-0.1048	0.1627	-0.4238	0.2141	-0.6442	0.521446	0.900	0.655	1.239
Previous tuberculosis	-0.0459	0.1928	-0.4239	0.3320	-0.2382	0.812374	0.955	0.655	1.394
Malnutrition	-0.1089	0.1895	-0.4803	0.2626	-0.5745	0.567370	0.897	0.619	1.300
Malignancy	0.1830	0.1666	-0.1434	0.5095	1.0990	0.275338	1.201	0.866	1.664
Methotrexate dosis	-0.0052	0.0094	-0.0237	0.0132	-0.5560	0.579875	0.995	0.977	1.013
Methotrexate time	-0.0010	0.0094	-0.0195	0.0175	-0.1040	0.917440	0.999	0.981	1.018
Leflunomide dosis	-0.0020	0.0086	-0.0189	0.0149	-0.2314	0.817620	0.998	0.981	1.015
Leflunomide time	-0.0169	0.0139	-0.0441	0.0103	-1.2171	0.227432	0.983	0.957	1.010
Prednisone dosis	0.0427	0.0176	0.0083	0.0772	2.4295	0.017546	1.044	1.008	1.080
Prednisone time	0.0122	0.0113	-0.0098	0.0343	1.0860	0.281003	1.012	0.990	1.035
Non-immunosuppressive prednisone	-0.0267	0.0242	-0.0741	0.0206	-1.1065	0.272094	0.974	0.929	1.021
Methotrexate + leflunomide dosis	0.0002	0.0006	-0.0010	0.0014	0.3128	0.755339	1.000	0.999	1.001
Methotrexate + prednisone dosis	-0.0031	0.0013	-0.0056	-0.0006	-2.3935	0.019224	0.997	0.994	0.999
Leflunomide + prednisone dosis	-0.0020	0.0014	-0.0047	0.0007	-1.4487	0.151647	0.998	0.995	1.001
Methotrexate + leflunomide + prednisone dosis	0.0001	0.0001	-0.0001	0.0003	1.2906	0.200868	1.000	1.000	1.000
Biological therapy	0.0757	0.0399	-0.0025	0.1540	1.8973	0.061693	1.079	0.998	1.166
Biological therapy + methotrexate dosis	0.0023	0.0018	-0.0014	0.0059	1.2185	0.226921	1.002	0.999	1.006
Biological therapy dosis	-0.0003	0.0003	-0.0010	0.0003	-1.0090	0.316254	1.000	0.999	1.000
Biological therapy time	-0.0351	0.0745	-0.1811	0.1109	-0.4714	0.638717	0.965	0.834	1.117

the development of anergy were evaluated through the logistic and Cox regression analysis for the dependent variables of anergy and non-anergy.

Ethical considerations

The present study meets the requirements for research in humans according to resolution 8430 of 1993 of the Ministry of Health. According to article 11 of the same resolution, the present study is classified as with minimal risk. It was approved by the ethics committee of the institutions involved.

Results

Patient characteristics

A total of 102 subjects were evaluated and 101 volunteers entered. One subject was excluded from the study because of an alternative diagnosis of osteoarthritis. Of the 101 volunteers, one was excluded due to confirmation of active tuberculosis in the follow up. There was no loss of information. [Figure 1](#) The median age of the participants was 57 years

(interquartile range = 12), 88% of these were women (n = 88). The time since diagnosis of RA was over 10 years in 42% of the patients (n = 42) and under 2 years in 15% (n = 15). The most frequent age group was 51 to 60 years old with 38% of the participants (n = 38). The most frequent comorbidity was diabetes mellitus in 9%. The most frequent immunosuppressive treatment was methotrexate followed by prednisone or equivalent and leflunomide with 74%, 69% and 46% of patients respectively. [Table 1](#) shows the summary of the sociodemographic characteristics of the volunteers.

Findings of skin tests

The frequency of cutaneous anergy was 9% (n = 9). It occurred in 33% of men versus 6% of women. Most cases presented the age group of 51 to 60 years with 56% (n = 5) and 89% were older than 50 years (n = 8). More than 10 years of illness was frequent with 44% of patients with anergy (n = 4). Only one of the patients with anergy presented associated comorbidity and it was due to diabetes. [Table 2](#). All patients with anergy were treated with corticosteroids (n = 9) and only one of these used low doses of prednisone or equivalent in monotherapy

Table 5 – Dependent variable logistic regression “Cutaneous energy”.

Category	Regression coefficients	Standard error	Lower limit (LL)	Upper limit (UL)	Critical value t	p Value	Odds Ratio (OR)	LL (OR)	UL(OR)
Intercept	0.1702	0.2735	-0.3658	0.7061	0.6223	0.535587			
Age	0.0010	0.0027	-0.0044	0.0063	0.3472	0.729436	1.001	0.996	1.006
Gender	-0.2300	0.0962	-0.4186	-0.0415	-2.3912	0.019266	0.795	0.658	0.959
Weight	0.0012	0.0028	-0.0043	0.0068	0.4364	0.663808	1.001	0.996	1.007
Years from diagnosis	0.0005	0.0027	-0.0047	0.0058	0.2037	0.839127	1.001	0.995	1.006
Diabetes	0.0444	0.1120	-0.1750	0.2638	0.3964	0.692901	1.045	0.839	1.302
Kidney disease	-0.1623	0.1624	-0.4807	0.1560	-0.9995	0.320733	0.850	0.618	1.169
Previous tuberculosis	-0.0633	0.2138	-0.4822	0.3557	-0.2960	0.768063	0.939	0.617	1.427
Malnutrition	-0.1980	0.2021	-0.5941	0.1981	-0.9796	0.330371	0.820	0.552	1.219
Malignancy	0.1523	0.1694	-0.1797	0.4844	0.8992	0.371384	1.165	0.836	1.623
Methotrexate dosis	-0.0054	0.0042	-0.0135	0.0027	-1.3005	0.197370	0.995	0.987	1.003
Methotrexate time	-0.0041	0.0099	-0.0235	0.0153	-0.4128	0.680928	0.996	0.977	1.015
Leflunomide dosis	-0.0013	0.0035	-0.0081	0.0056	-0.3594	0.720329	0.999	0.992	1.006
Leflunomide time	-0.0194	0.0140	-0.0469	0.0082	-1.3777	0.172332	0.981	0.954	1.008
Antimalarial dosis	0.0008	0.0006	-0.0005	0.0021	1.2624	0.210672	1.001	1.000	1.002
Antimalarial time	-0.0215	0.0293	-0.0789	0.0359	-0.7336	0.465449	0.979	0.924	1.037
Sulfasalazine dosis	-0.0001	0.0001	-0.0003	0.0001	-0.8457	0.400351	1.000	1.000	1.000
Sulfasalazine time	-0.0103	0.0310	-0.0709	0.0504	-0.3322	0.740657	0.990	0.932	1.052
Prednisone dosis	0.0115	0.0102	-0.0085	0.0316	1.1269	0.263350	1.012	0.992	1.032
Prednisone time	0.0090	0.0117	-0.0138	0.0319	0.7740	0.441327	1.009	0.986	1.032
Biological therapy	0.0914	0.0386	0.0158	0.1670	2.3689	0.020382	1.096	1.016	1.182
Biological therapy dosis	-0.0002	0.0003	-0.0009	0.0004	-0.6841	0.495988	1.000	0.999	1.000
Biological therapy time	-0.0263	0.0931	-0.2089	0.1562	-0.2829	0.778048	0.974	0.811	1.169

(less than or equal to 5 mg), methotrexate was used in 44% (n = 4) and biological therapy in 33% (n = 3).

The overall positivity for tuberculin was 13%. Of these, 62% were asymptomatic (n = 8) and 38% were symptomatic (n = 5). The most frequent symptom was unexplained weight loss in 31% (n = 4) followed by cough and sweating with 23% each (n = 3). Only 15% of patients experienced fever (n = 2). None of the patients experienced dyspnea or hemoptysis.

When assessing whether anergy presence is influenced by other variables, a multivariate analysis was performed with the logistic regression model. A model was designed in which the variables considered relevant were included. This model was evaluated for the presence of interactions (Comorbidities, sociodemographic variables, dose and interaction time of immunosuppressive medications). [Table 3](#). In this model, female sex behaved as a protective variable for the generation of anergy OR 0.795 [95% CI, 0.658 - 0.959, p <0.05] and biological therapy as a risk factor OR 1.096 [95% CI, 1.016-1.182, p <0.05]. No associations with statistical significance were found in the rest of the analyzed variables.

Another logistic regression model was constructed by combining variables and distinguishing between immunosuppressive versus non-immunosuppressive doses of prednisone (less than or equal to 5 mg in monotherapy). [Table 4](#). No statistical significance was found in the combination of immunosuppressive treatments. Treatment with prednisone in immunosuppressive doses is associated as a risk factor for the presentation of anergy OR 1,044 [95% CI, 1,008-1080 p <0.05]. [Table 5](#)

Discussion

RA is an autoimmune, inflammatory, chronic and progressive disease, characterized mainly by the damage of small joints

of the hands and feet. It is recommended that all patients diagnosed with RA begin with therapy with disease-modifying antirheumatic drugs (DMARDs) and to a large extent many require scaling up to biological therapy.^{73,74} RA per se has been associated with the reactivation of latent TB.¹ There are multiple studies that show a decrease in tuberculin positivity in patients with RA.^{18,66,75-77} Immunosuppressive drug therapy, particularly iTNF, significantly increases this risk.^{71,78,79}

Routine use of hypersensitivity tests is recommended for the diagnosis of latent TB and thus prevent reactivation. There are data that suggest that the use of a single screening test does not identify all patients at risk of TB, since false negative results are more likely in immunocompromised individuals.^{80,81} The dual test strategy (tuberculin + IGRA) or the use of a tuberculin booster is consistent with the recommendations of the American College of Rheumatology and other public health agencies.^{71,78,82}

Some reports consider that the use of disease-modifying therapy and steroids are a cause of anergy without this being proven by comparison to cutaneous controls or serological methods for the diagnosis of TB.^{67,77,83} This is the first study that evaluates the prevalence of cutaneous anergy in patients with RA and seeks to determine if there are variables associated with its appearance. [Table 6](#)

Anergy is defined as the non-induration of any intradermal antigen whose immune response is of high prevalence in the general population. In this study, the tetanus toxoid concomitant with the application of tuberculin was chosen.

We found that a significant proportion of patients had anergy in up to 1 in 11 patients (9%). This is consistent with the recommendation of experts to use a combination of tests that would increase sensitivity.^{18,71}

The main risk factors associated with cutaneous anergy in various studies are acquired immunosuppression states such

Table 6 – Logistic regression combined variables.

Category	Regression coefficients	Standard error	Lower limit (LL)	Upper limit (UL)	Critical value t	p Value	Odds Ratio (OR)	LL (OR)	UL(OR)
Intercept	0.3903	0.5484	-0.6845	1.4651	0.7117	0.478858			
Age	0.0012	0.0026	-0.0039	0.0063	0.4653	0.643053	1.001	0.996	1.006
Gender	-0.2111	0.0899	-0.3873	-0.0349	-2.3484	0.021528	0.810	0.679	0.966
Weight	0.0004	0.0028	-0.0051	0.0059	0.1515	0.880004	1.000	0.995	1.006
Years from diagnosis	0.0007	0.0028	-0.0047	0.0061	0.2459	0.806429	1.001	0.995	1.006
Diabetes	0.0768	0.1076	-0.1341	0.2878	0.7138	0.477615	1.080	0.874	1.333
Kidney disease	-0.1048	0.1627	-0.4238	0.2141	-0.6442	0.521446	0.900	0.655	1.239
Previous tuberculosis	-0.0459	0.1928	-0.4239	0.3320	-0.2382	0.812374	0.955	0.655	1.394
Malnutrition	-0.1089	0.1895	-0.4803	0.2626	-0.5745	0.567370	0.897	0.619	1.300
Malignancy	0.1830	0.1666	-0.1434	0.5095	1.0990	0.275338	1.201	0.866	1.664
Methotrexate dosis	-0.0052	0.0094	-0.0237	0.0132	-0.5560	0.579875	0.995	0.977	1.013
Methotrexate time	-0.0010	0.0094	-0.0195	0.0175	-0.1040	0.917440	0.999	0.981	1.018
Leflunomide dosis	-0.0020	0.0086	-0.0189	0.0149	-0.2314	0.817620	0.998	0.981	1.015
Leflunomide time	-0.0169	0.0139	-0.0441	0.0103	-1.2171	0.227432	0.983	0.957	1.010
Prednisone dosis	0.0427	0.0176	0.0083	0.0772	2.4295	0.017546	1.044	1.008	1.080
Prednisone time	0.0122	0.0113	-0.0098	0.0343	1.0860	0.281003	1.012	0.990	1.035
Non-immunosuppressive prednisone	-0.0267	0.0242	-0.0741	0.0206	-1.1065	0.272094	0.974	0.929	1.021
Methotrexate + leflunomide dosis	0.0002	0.0006	-0.0010	0.0014	0.3128	0.755339	1.000	0.999	1.001
Methotrexate + prednisone dosis	-0.0031	0.0013	-0.0056	-0.0006	-2.3935	0.019224	0.997	0.994	0.999
Leflunomide + prednisone dosis	-0.0020	0.0014	-0.0047	0.0007	-1.4487	0.151647	0.998	0.995	1.001
Methotrexate + leflunomide + prednisone dosis	0.0001	0.0001	-0.0001	0.0003	1.2906	0.200868	1.000	1.000	1.000
Biological therapy	0.0757	0.0399	-0.0025	0.1540	1.8973	0.061693	1.079	0.998	1.166
Biological therapy + methotrexate dosis	0.0023	0.0018	-0.0014	0.0059	1.2185	0.226921	1.002	0.999	1.006
Biological therapy dosis	-0.0003	0.0003	-0.0010	0.0003	-1.0090	0.316254	1.000	0.999	1.000
Biological therapy time	-0.0351	0.0745	-0.1811	0.1109	-0.4714	0.638717	0.965	0.834	1.117

as malnutrition, hematological tumors and HIV patients with low CD4 counts.^{37,39,51,58,71,84-86} In other diseases that receive a similar degree of immunosuppression to RA, the prevalence of anergy can reach 83%.⁶⁸ In rheumatologic diseases, there are few published studies although there is clear evidence of alteration of cellular immunity.^{87,88} In a study conducted by Ponce de León, in patients with RA the frequency of negative PPD was 70% vs 26% in healthy controls.⁸⁹

In the present study it is striking that all subjects with anergy used corticosteroids. This variable significantly increases the probability of presenting anergy and is dose dependent, presenting with doses greater than 5 mg or in combination therapy. In contrast, and despite the biological plausibility, no association was found in the use of DMARDs, particularly metotrexate, the most frequent DMARD agent; as found in other studies in animals and immunosuppressed patients.⁹⁰⁻⁹³ In favor of this finding, there are studies in which a paradoxical effect of metotrexate has been found with an increase in false positives of PPD.⁹⁴

As expected, we found that the use of biological therapy is a significant risk for the generation of anergy. This group is of special importance since there are no recommendations

about screening or diagnostic studies of latent TB in patients who are already on biologic therapy.

In the present study, a third of the patients with anergy were older than 60 years. There are studies that report that older subjects may have lower antigenic reactivity.^{87,88,95} However, we did not find that age was a determining factor of anergy in patients. There was no anergy in subjects older than 80 years and the most prevalent group was 51 to 60 years.

The female gender behaved as a protective factor to present anergy, there's plausibility that sex hormones have a role in the cellular immune response that needs further research.⁹⁶

The prevalence of positive tuberculin in patients with RA was low in this study (13%) compared to other studies in high prevalence countries where it ranges between 20 and 40%.^{66,94,97} In the Indian study by Agarwal et al.⁶⁶ a prevalence of tuberculin positivity of 20.4% was found. Similarly to the present study, it was found that the use of steroids decreased the reaction to tuberculin (3% versus 25%, $P = 0.002$). They also found no association with the use of other DMARDs.

Our study has several limitations. First, the small number of subjects ($n = 100$) can reduce the chances of obtaining statistically significant results. Additionally, the prevalence of anergy

was lower than expected, which compromises the validity of the association measures. Thirdly, we did not analyze the correlation between anergy results and disease activity.

Conclusion

The high prevalence of cutaneous anergy in patients with RA in the present study and the evidence presented here supports the recommendation of a second diagnostic test (tuberculin booster or IGRAs) for the diagnosis of latent TB in patients with RA and immunosuppressive therapy.

Funding

This work was funded through an internal call from the National University of Colombia “national call for projects to strengthen the research, creation and innovation of the national university of Colombia 2016-2018” The project identifies with the Hermes code: 37595 and QUIPU code: 201010028367. It also received financial support from the Colombian Association of Rheumatology.

Conflict interest

The authors declare that they have no financial or personal relationship with people or organizations that could give rise to a conflict of interest in relation to the present study.

REFERENCES

- Carmona L, Hernández-García C, Vadillo C, Pato E, Balsa A, González-Alvaro I, et al. Increased risk of tuberculosis in patients with rheumatoid arthritis. *J Rheumatol*. 2003;30:1436-9.
- Dixon WG, Hyrich KL, Watson KD, Lunt M, Galloway J, Ustianowski A, et al. Drug-specific risk of tuberculosis in patients with rheumatoid arthritis treated with anti-TNF therapy: results from the British Society for Rheumatology Biologics Register (BSRBR). *Ann Rheum Dis*. 2010;69:522-8, <http://dx.doi.org/10.1136/ard.2009.118935>.
- Mack U, Migliori GB, Sester M, Rieder HL, Ehlers S, Goletti D, et al. LTBI: latent tuberculosis infection or lasting immune responses to M. tuberculosis? A TBNET consensus statement. *Eur Respir J*. 2009;33:956-73, <http://dx.doi.org/10.1183/09031936.00120908>.
- Targeted tuberculin testing and treatment of latent tuberculosis infection. American Thoracic Society. *MMWR Recomm Rep*. 2000 9;49(RR-6):1-51.
- Getahun H, Matteelli A, Chaisson RE, Raviglione M. Latent mycobacterium tuberculosis infection. *N Engl J Med*. 2015;372:2127-35, <http://dx.doi.org/10.1056/NEJMra1405427>.
- Corbett EL, Watt CJ, Walker N, Maher D, Williams BG, Raviglione MC, et al. The growing burden of tuberculosis: global trends and interactions with the HIV epidemic. *Arch Intern Med*. 2003;163:1009-21, <http://dx.doi.org/10.1001/archinte.163.9.1009>.
- Comstock GW, Livesay VT, Woolpert SF. The prognosis of a positive tuberculin reaction in childhood and adolescence. *Am J Epidemiol*. 1974;99:131-8, <http://dx.doi.org/10.1093/oxfordjournals.aje.a121593>.
- Jick SS, Lieberman ES, Rahman MU, Choi HK. Glucocorticoid use, other associated factors, and the risk of tuberculosis. *Arthritis Rheum*. 2006;55:19-26, <http://dx.doi.org/10.1002/art.21705>.
- Keane J, Bresnahan B. Tuberculosis reactivation during immunosuppressive therapy in rheumatic diseases: diagnostic and therapeutic strategies. *Curr Opin Rheumatol*. 2008;20:443-9, <http://dx.doi.org/10.1097/BOR.0b013e3283025ec2>.
- American College of Rheumatology Subcommittee on Rheumatoid Arthritis Guidelines. Guidelines for the management of rheumatoid arthritis: 2002 update. *Arthritis Rheum*. 2002;46:328-46, <https://doi.org/10.1002/art.10148>.
- Day R. Adverse reactions to TNF- α inhibitors in rheumatoid arthritis. *The Lancet*. 2002;359:540-1, [http://dx.doi.org/10.1016/s0140-6736\(02\)07718-8](http://dx.doi.org/10.1016/s0140-6736(02)07718-8).
- Chen C, Raisch DW. Post-marketing research studies of the effectiveness and safety of biologics for rheumatoid arthritis treatment: a systematic literature review. *Value in Health*. 2016;19:A224-5, <http://dx.doi.org/10.1016/j.jval.2016.03.1159>.
- Latent tuberculosis infection: updated and consolidated guidelines for programmatic management. Geneva: World Health Organization; 2018.
- Allen A, Carville S, McKenna F, Guideline Development Group. Diagnosis and management of rheumatoid arthritis in adults: summary of updated NICE guidance. *BMJ*. 2018;362:k3015, <http://dx.doi.org/10.1136/bmj.k3015>.
- Gómez Reino J, Loza E, Andreu JL, Balsa A, Batlle E, Cañete JD, et al. [Consensus statement of the Spanish Society of Rheumatology on risk management of biologic therapy in rheumatic patients]. *Reumatol Clin*. 2011;7:284-98, <http://dx.doi.org/10.1016/j.reuma.05.002.2011>.
- Bombardier C, Hazlewood GS, Akhavan P, Schieir O, Dooley A, Haraoui B, et al. Canadian Rheumatology Association recommendations for the pharmacological management of rheumatoid arthritis with traditional and biologic disease-modifying antirheumatic drugs: part II safety. *J Rheumatol*. 2012;39:1583-602, <http://dx.doi.org/10.3899/jrheum.65.1201>.
- Dogan E, Erkok R, Sayarlioglu H, Uzun K. Tuberculin skin test results and the booster phenomenon in two-step tuberculin skin testing in hemodialysis patients. *Ren Fail*. 2005;27:425-8.
- Pérez-Barbosa L, Esquivel-Valerio JA, Arana-Guajardo AC, Vega-Morales D, Riega-Torres J, Garza-Elizondo MA. Increased detection of latent tuberculosis by tuberculin skin test and booster phenomenon in early rheumatoid arthritis patients. *Rheumatol Int*. 2015;35:1555-9, <http://dx.doi.org/10.1007/s00296-015-93246>.
- Smolen JS, Landewé R, Bijlsma J, Burmester G, Chatzidionysiou K, Dougados M, et al. EULAR recommendations for the management of rheumatoid arthritis with synthetic and biological disease-modifying antirheumatic drugs: 2016 update. *Ann Rheum Dis*. 2017;76:960-77, <http://dx.doi.org/10.1136/annrheumdis-2016-210715>.
- Trajman A, Steffen RE, Menzies D. Interferon-gamma release assays versus tuberculin skin testing for the diagnosis of latent tuberculosis infection: an overview of the evidence. *Pulm Med*. 2013;2013:601737, <http://dx.doi.org/10.1155/2013/601737>.
- Nienhaus A, Schablon A, Diel R. Interferon-gamma release assay for the diagnosis of latent tb infection - Analysis of discordant results, when compared to the tuberculin skin test. *PLoS One*. 2008;3:e2665, <http://dx.doi.org/10.1371/journal.pone.0002665>.
- Diagnostic standards and classification of tuberculosis in adults and children. This official statement of the American Thoracic Society and the Centers for Disease Control and

- Prevention was adopted by the ATS Board of Directors, July 1999. This statement was endorsed by the Council of the Infectious Disease Society of America, September 1999. *Am J Respir Crit Care Med.* 2000; 161(4 Pt 1):1376-95. 10.1164/ajrccm.161.4.16141.
23. Dheda K, Schwander SK, Zhu B, van Zyl-Smit RN, Zhang Y. The immunology of tuberculosis: from bench to bedside. *Respirology.* 2010;15:433-50, <http://dx.doi.org/10.1111/j.1440-1843.2010.01739.x>.
 24. Tsicopoulos A, Hamid Q, Varney V, Ying S, Moqbel R, Durham SR, et al. Preferential messenger RNA expression of Th1-type cells (IFN-gamma+ IL-2+) in classical delayed-type (tuberculin) hypersensitivity reactions in human skin. *J Immunol.* 1992;148:2058-61.
 25. Sokal JE. Measurement of delayed skin-test responses. *N Engl J Med.* 1975;293:501-2, <http://dx.doi.org/10.1056/NEJM197509042931013>.
 26. Jordan TJ, Sunderam G, Thomas L, Reichman LB. Tuberculin reaction size measurement by the pen method compared to traditional palpation *Chest.* 1987;92:234-6, <http://dx.doi.org/10.1378/chest.92.2.234>.
 27. Shankar MSR, Ravi Shankar MS, Aravindan AN, Sohal PM, Kohli HS, Sud K, et al. The prevalence of tuberculin sensitivity and anergy in chronic renal failure in an endemic area: tuberculin test and the risk of post-transplant tuberculosis. *Nephrol Dial Transplant.* 2005;20:2720-4, <http://dx.doi.org/10.1093/ndt/gfi141>.
 28. Poduval RD, Hammes MS. Tuberculosis screening in dialysis patients- is the tuberculin test effective? *Clin Nephrol.* 2003;59:436-40, <http://dx.doi.org/10.5414/cnp59436>.
 29. Smirnoff M, Patt C, Seckler B, Adler JJ. Tuberculin and anergy skin testing of patients receiving long-term hemodialysis. [Internet]. *Chest.* 1998;113:25-7, <http://dx.doi.org/10.1378/chest.113.1.25>.
 30. Woeltje KF, Mathew A, Rothstein M, Seiler S, Fraser VJ. Tuberculosis infection and anergy in hemodialysis patients. *Am J Kidney Dis.* 1998;31:848-52, [http://dx.doi.org/10.1016/s0272-6386\(98\)70055-1](http://dx.doi.org/10.1016/s0272-6386(98)70055-1).
 31. Pesanti EL. The negative tuberculin test. Tuberculin, HIV, and anergy panels. *Am J Respir Crit Care Med.* 1994;149:1699-709, <http://dx.doi.org/10.1164/ajrccm.149.6.7710481>.
 32. García-García ML, Valdespino-Gómez JL, García-Sancho C, Mayar-Maya ME, Palacios-Martínez M, Balandrano-Campos S, et al. Underestimation of Mycobacterium tuberculosis infection in HIV-infected subjects using reactivity to tuberculin and anergy panel. *Int J Epidemiol.* 2000;29:369-75, <http://dx.doi.org/10.1093/ije/29.2.369>.
 33. Daniel TM, Janicki BW. Mycobacterial antigens: a review of their isolation, chemistry, and immunological properties. *Microbiol Rev.* 1978;42:84-113.
 34. Harboe M. Antigens of PPD, old tuberculin, and autoclaved Mycobacterium bovis BCG studied by crossed immunoelectrophoresis. *Am Rev Respir Dis.* 1981;124:80-7, <http://dx.doi.org/10.1164/arrd.1981.124.1.80>.
 35. Von Pirquet CE. Allergy. *Arch Intern Med (Chic).* 1911;VII(2):259-88. 10.1001/archinte.1911.00060020128010.
 36. Colebunders RL, Lebughe I, Nzila N, Kalunga D, Francis H, Ryder R, et al. Cutaneous delayed-type hypersensitivity in patients with human immunodeficiency virus infection in Zaire. *J Acquir Immune Defic Syndr.* 1989;2:576-8.
 37. Graham NM, Nelson KE, Solomon L, Bonds M, Rizzo RT, Scavotto J, et al. Prevalence of tuberculin positivity and skin test anergy in HIV-1-seropositive and -seronegative intravenous drug users. *JAMA.* 1992;267:369-73.
 38. Pulmonary Complications of HIV Infection Study Group, Markowitz N, Hansen NI, Wilcosky TC, Hopewell PC, Glassroth J, Kvale PA, et al. Tuberculin and anergy testing in HIV-seropositive and HIV-seronegative persons. *Ann Intern Med.* 1993;119:185-93, <http://dx.doi.org/10.7326/0003-4819-119-3-199308010-20000>.
 39. Pelly TF, Santillan CF, Gilman RH, Cabrera LZ, Garcia E, Vidal C, et al. Tuberculosis skin testing, anergy and protein malnutrition in Peru. *Int J Tuberc Lung Dis.* 2005;9:977-84.
 40. Bansal VK, Popli S, Pickering J, Ing TS, Vertuno LL, Hano JE. Protein-calorie malnutrition and cutaneous anergy in hemodialysis maintained patients. *Am J Clin Nutr.* 1980;33:1608-11, <http://dx.doi.org/10.1093/ajcn/33.7.1608>.
 41. Boussiotis VA, Tsai EY, Yunis EJ, Thim S, Delgado JC, Dascher CC, et al. IL-10-producing T cells suppress immune responses in anergic tuberculosis patients. *J Clin Invest.* 2000;105:1317-25, <http://dx.doi.org/10.1172/JCI9918>.
 42. Delafuente JC, Eisenberg JD, Hoelzer DR, Slavin RG. Tetanus toxoid as an antigen for delayed cutaneous hypersensitivity. *JAMA.* 1983;249:3209-11.
 43. Whittingham S, Feery B, Mackay IR. Use of tetanus toxoid for testing cell-mediated immunity. *Aust N Z J Med.* 1982;12:511-4, <http://dx.doi.org/10.1111/j.1445-5994.1982.tb03833.x>.
 44. Salesi M, Meidani M, Meshkinfar S, Hashemi H, Farajzadegan Z. Purified protein derivative test and its booster phenomenon in patients with rheumatoid arthritis. *Adv Biomed Res.* 2015;4:80, <http://dx.doi.org/10.4103/2277-9175.156638>.
 45. Bathon JM, Cohen SB. The 2008 American College of Rheumatology recommendations for the use of nonbiologic and biologic disease-modifying antirheumatic drugs in rheumatoid arthritis: Where the rubber meets the road *Arthritis Rheum.* 2008;59:757-9, <http://dx.doi.org/10.1002/art.23723>.
 46. Sokal JE, Primikiri N. The delayed skin test response in Hodgkin's disease and lymphosarcoma Effect of disease activity. *Cancer.* 1961;14:597-607, [http://dx.doi.org/10.1002/1097-0142\(199005/06\)14:3<597::aid-cnrcr2820140321>3.0.co;2-x](http://dx.doi.org/10.1002/1097-0142(199005/06)14:3<597::aid-cnrcr2820140321>3.0.co;2-x).
 47. Kataria YP, Sagone AL, LoBuglio AG, Bromberg PA. In vitro observations on sarcoid lymphocytes and their correlation with cutaneous energy and clinical severity of disease. *Am Rev Respir Dis.* 1973;108:767-76, <http://dx.doi.org/10.1164/arrd.1973.108.4.767>.
 48. Friou GJ. A study of the cutaneous reactions to oidiomycin, trichophytin, and mumps skin test antigens in patients with sarcoidosis. *Yale J Biol Med.* 1952;24:533-9.
 49. Bratt G, von Krogh G, Moberg L, Karlsson A, Putkonen PO, Biberfeld G, et al. Intradermal testing with multiple recall antigens for identification of cell-mediated immune deficiency in homosexual men. *Clin Immunol Immunopathol.* 1986;41:206-15, [http://dx.doi.org/10.1016/0090-1229\(86\)90104-2](http://dx.doi.org/10.1016/0090-1229(86)90104-2).
 50. Sears SD, Fox R, Brookmeyer R, Leavitt R, Frank Polk B. Delayed hypersensitivity skin testing and anergy in a population of gay men *Clin Immunol Immunopathol.* 1987;45:177-83, [http://dx.doi.org/10.1016/0090-1229\(87\)90032-8](http://dx.doi.org/10.1016/0090-1229(87)90032-8).
 51. Schier WW. Cutaneous anergy and Hodgkin's disease. *N Engl J Med.* 1954;250:353-61, <http://dx.doi.org/10.1056/NEJM195403042500902>.
 52. Altered reactivity to skin homografts in severe thermal injury. *Plastic and Reconstructive Surgery.* 1964;34(2):216-7, <http://dx.doi.org/10.1097/00006534-196408000-00035>.
 53. Facktor M, Bernstein R, Fireman P. Hypersensitivity to tetanus toxoid. *J Allergy Clin Immunol.* 1973;52:1-12, [http://dx.doi.org/10.1016/0091-6749\(73\)90115-2](http://dx.doi.org/10.1016/0091-6749(73)90115-2).
 54. Callaghan JT, Petersen BH, Smith WC, Epinette WW, Ransburg RC. Delayed hypersensitivity to mumps antigen in humans *Clin Immunol Immunopathol.* 1983;26:102-10, [http://dx.doi.org/10.1016/0090-1229\(83\)90178-2](http://dx.doi.org/10.1016/0090-1229(83)90178-2).
 55. Selwyn PA, Sckell BM, Alcabes P, Friedland GH, Klein RS, Schoenbaum EE. High risk of active tuberculosis in

- HIV-infected drug users with cutaneous anergy. *JAMA*. 1992;268:504-9.
56. Dale JB. Type-specific immunogenicity of a chemically synthesized peptide fragment of type 5 streptococcal M protein. *J Exp Med*. 1983;158:1727-32, <http://dx.doi.org/10.1084/jem.158.5.1727>.
 57. Galant SP, Flod N, Shimizu I, Granger GA, Groncy CE. Relationship between cutaneous delayed hypersensitivity and cell-mediated immunity in vitro responses assessed by diphtheria and tetanus toxoids. *J Allergy Clin Immunol*. 1977;60:247-53, [http://dx.doi.org/10.1016/0091-6749\(77\)90139-7](http://dx.doi.org/10.1016/0091-6749(77)90139-7).
 58. The Pulmonary Complications of HIV Infection Study Group, Chin DP, Osmond D, Page-Shafer K, Glassroth J, Rosen MJ, Reichman LB, et al. Reliability of anergy skin testing in persons with HIV infection. *Am J Respir Crit Care Med*. 1996;153 Pt 1:1982-4, <http://dx.doi.org/10.1164/ajrccm.153.6.8665065>.
 59. Turk JL, Rudner EJ, Heather CJ. A histochemical analysis of mononuclear cell infiltrates of the skin II. Delayed hypersensitivity in the human. *Int Arch Allergy Appl Immunol*. 1966;30:248-56, <http://dx.doi.org/10.1159/000229810>.
 60. Gell PGH, Hinde IT. The histology of the tuberculin reaction and its modification by cortisone. *Br J Exp Pathol*. 1951;32:516-29.
 61. Mccluskey RT, Benacerraf B, Mccluskey JW. Studies on the specificity of the cellular infiltrate in delayed hypersensitivity reactions. *J Immunol*. 1963;90:466-77.
 62. Johnson C, Walls RS, Ruwoldt A. Delayed hypersensitivity to tetanus toxoid in man: in vivo and in vitro studies. *Pathology*. 1983;15:369-72, <http://dx.doi.org/10.3109/00313028309085161>.
 63. Borut TC, Ank BJ, Richard Stiehm E. Tetanus skin test: does it assess delayed hypersensitivity? *Pediatric Research*. [Internet]. 1977;11:485, <http://dx.doi.org/10.1203/00006450-197704000-00689>.
 64. French AL, McCullough ME, Rice KT, Schultz ME, Gordin FM. The use of tetanus toxoid to elucidate the delayed-type hypersensitivity response in an older, immunized population. *Gerontology*. 1998;44:56-60, <http://dx.doi.org/10.1159/000021984>.
 65. Stein M, Sela-Razon B, Kleter Y, Somekh E. Reliability of control skin tests with common antigens in children undergoing tuberculin skin test. *Ann N Y Acad Sci*. 2007;1109:235-9, <http://dx.doi.org/10.1196/annals.1398.028>.
 66. Agarwal S, Das SK, Agarwal GG, Srivastava R. Steroids decrease prevalence of positive tuberculin skin test in rheumatoid arthritis: implications on anti-TNF therapies. *Interdiscip Perspect Infect Dis*. 2014;2014:430134, <http://dx.doi.org/10.1155/2014/430134>.
 67. Winkelstein A. Effects of cytotoxic immunosuppressants on tuberculin-sensitive lymphocytes in guinea pigs. *J Clin Invest*. 1975;56:1587-96, <http://dx.doi.org/10.1172/JCI108241>.
 68. Mow WS, Abreu-Martin MT, Papadakis KA, Pitchon HE, Targan SR, Vasiliauskas EA. High incidence of anergy in inflammatory bowel disease patients limits the usefulness of PPD screening before infliximab therapy. *Clin Gastroenterol Hepatol*. 2004;2:309-13, [http://dx.doi.org/10.1016/s1542-3565\(04\)00060-6](http://dx.doi.org/10.1016/s1542-3565(04)00060-6).
 69. Stimpson PG, Paty JG Jr, Hudson T, Lieberman P. Delayed hypersensitivity skin testing for assessing anergy in the mid-south. *South Med J*. 1976;69:424-6, <http://dx.doi.org/10.1097/00007611-197604000-00013>.
 70. Kniker WT, Anderson CT, McBryde JL, Roumiantzeff M, Lesourd B, Multitest CMI. for standardized measurement of delayed cutaneous hypersensitivity and cell-mediated immunity Normal values and proposed scoring system for healthy adults in the U.S.A. *Ann Allergy*. 1984;52:75-82.
 71. Lewinsohn DM, Leonard MK, LoBue PA, Cohn DL, Daley CL, Desmond E, et al. Official American Thoracic Society/Infectious Diseases Society of America/Centers for Disease Control and Prevention Clinical Practice Guidelines: Diagnosis of tuberculosis in adults and children. *Clin Infect Dis*. 2017;64:111-5, <http://dx.doi.org/10.1093/cid/ciw694>.
 72. Palinkas LA, Horwitz SM, Green CA, Wisdom JP, Duan N, Hoagwood K. Purposeful Sampling for qualitative data collection and analysis in mixed method implementation research. *Adm Policy Ment Health*. 2015;42:533-44, <http://dx.doi.org/10.1007/s10488-013-0528-y>.
 73. Singh JA, Furst DE, Bharat A, Curtis JR, Kavanaugh AF, Kremer JM, et al. 2012 update of the 2008 American College of Rheumatology recommendations for the use of disease-modifying antirheumatic drugs and biologic agents in the treatment of rheumatoid arthritis. *Arthritis Care Res (Hoboken)*. 2012;64:625, <http://dx.doi.org/10.1002/acr.21641>.
 74. Smolen JS, Breedveld FC, Burmester GR, Bykerk V, Dougados M, Emery P, et al. Treating rheumatoid arthritis to target: 2014 update of the recommendations of an international task force. *Ann Rheum Dis*. 2016;75:3-15, <http://dx.doi.org/10.1136/annrheumdis-2015-207524>.
 75. Marques CDL, Duarte ÂLBP, Lorena VMB de, Souza JR de, Souza W, Gomes Y de M, et al. Resposta atenuada ao PPD no diagnóstico de infecção tuberculosa latente em pacientes com artrite reumatoide. *Rev Bras Reumatol*. 2009; 49(2):121-5.
 76. Arredondo AM, Escobar-Trujillo A, Londono J, Gonzalez-Malaver F, Bello-Gualtero J, Guzman-Vergara C, et al. AB0203 prevalence of tuberculin skin test reaction positivity in a Colombian cohort of rheumatoid arthritis patients on biologic therapy [Internet]. *Ann Rheum Diseases*. 2013;72:A48.
 77. Karkucak M, Capkin E, Ozsu S, Nuhoglu I, Erol M, Yilmaz G, et al. An evaluation of the tuberculin skin test for anti TNF alpha prophylaxis in patients with ankylosing spondylitis and rheumatoid arthritis. *Bratisl Lek Listy*. 2010;111:498-501.
 78. Calabrese LH, Calabrese C, Kirchner E. The 2015 American College of Rheumatology Guideline for the Treatment of Rheumatoid Arthritis should include new standards for hepatitis b screening: comment on the article by Singh et al. *Arthritis Rheumatol*. 2016;68:1314-5, <http://dx.doi.org/10.1002/art.39635>.
 79. Gómez-Reino JJ, Carmona L, Valverde VR, Mola EM, Montero MD. Treatment of rheumatoid arthritis with tumor necrosis factor inhibitors may predispose to significant increase in tuberculosis risk: A multicenter active-surveillance report. *Arthritis Rheum*. 2003;48:2122-7, <http://dx.doi.org/10.1002/art.11137>.
 80. Kleinert S, Tony H-P, Krueger K, Detert J, Mielke F, Rockwitz K, et al. Screening for latent tuberculosis infection: performance of tuberculin skin test and interferon- γ release assays under real-life conditions. *Ann Rheum Dis*. 2012;71:1791-5, <http://dx.doi.org/10.1136/annrheumdis-2011-200941>.
 81. Mariette X, Baron G, Tubach F, Lioté F, Combe B, Miceli-Richard C, et al. Influence of replacing tuberculin skin test with ex vivo interferon γ release assays on decision to administer prophylactic antituberculosis antibiotics before anti-TNF therapy. *Ann Rheum Dis*. 2012;71:1783-90, <http://dx.doi.org/10.1136/annrheumdis-2011-200408>.
 82. Winthrop KL, Weinblatt ME, Daley CL. You can't always get what you want, but if you try sometimes (with two tests —TST and IGRA— for tuberculosis) you get what you need. *Ann Rheum Dis*. 2012;71:1757-60, <http://dx.doi.org/10.1136/annrheumdis-2012-201979>.
 83. Iannone F, Cantini F, Lapadula G. Diagnosis of latent tuberculosis and prevention of reactivation in rheumatic patients receiving biologic therapy: international recommendations. *J Rheumatol Suppl*. 2014;91:41-6, <http://dx.doi.org/10.3899/jrheum.140101>.

84. Yanai H, Uthairavit W, Mastro TD, Limpakarnjanarat K, Sawanpanyalert P, Morrow RH Jr, et al. Utility of tuberculin and anergy skin testing in predicting tuberculosis infection in human immunodeficiency virus-infected persons in Thailand. *Int J Tuberc Lung Dis.* 1997;1:427-34.
85. Schier WW. Cutaneous anergy and Hodgkin's disease. *N Engl J Med.* 1954;250:353-61, <http://dx.doi.org/10.1056/NEJM195403042500902>.
86. Graham NMH. Prevalence of tuberculin positivity and skin test anergy in HIV-1—seropositive and —seronegative intravenous drug users. *JAMA.* 1992;267:369, <http://dx.doi.org/10.1001/jama.1992.03480030047035>.
87. Dorken E, Grzybowski S, Allen EA. Significance of the tuberculin test in the elderly. *Chest.* 1987;92:237-40, <http://dx.doi.org/10.1378/chest.92.2.237>.
88. Panayi GS, Corrigan VM, Pitzalis C. Pathogenesis of rheumatoid arthritis The role of T cells and other beasts. *Rheum Dis Clin North Am.* 2001;27:317-34, [http://dx.doi.org/10.1016/s0889-857x\(05\)70204-0](http://dx.doi.org/10.1016/s0889-857x(05)70204-0).
89. Ponce de León D, Acevedo-Vásquez E, Sánchez-Torres A, Cucho M, Alfaro J, Perich R, et al. Attenuated response to purified protein derivative in patients with rheumatoid arthritis: study in a population with a high prevalence of tuberculosis *Ann Rheum Dis.* 2005;64:1360-1, <http://dx.doi.org/10.1136/ard.2004.029041>.
90. Friedman RM, Buckler CE. Methotrexate inhibition of tuberculin hypersensitivity in inbred Guinea pigs. *J Immunol.* 1963;91:846-50.
91. Friedman RM. Inhibition of established tuberculin hypersensitivity by methotrexate *Proc Soc Exp Biol Med.* 1964;116:471-5, <http://dx.doi.org/10.3181/00379727-116-29282>.
92. Hart PD, Rees RJ, Niven JS. The effect of high dosage of methotrexate, associated with folinic acid, on the suppression of tuberculin sensitivity in Guinea-pigs. *Clin Exp Immunol.* 1968;3:91-8.
93. Ravindran V. From the editor's desk. *Indian J Rheumatol.* 2017;12:1.
94. Arias-Guillén M, Sánchez Menéndez MM, Alperi M, Riestra S, González Budiño MT, García-Clemente MM, et al. High rates of tuberculin skin test positivity due to methotrexate therapy: False positive results? *Semin Arthritis Rheum.* 2018;48:538-46, <http://dx.doi.org/10.1016/j.semarthrit.2018.03.018>.
95. Delafuente JC. Tetanus toxoid as an antigen for delayed cutaneous hypersensitivity. *JAMA.* 1983;249:3209, <http://dx.doi.org/10.1001/jama.1983.03330470049031>.
96. Klein SL, Flanagan KL. Sex differences in immune responses *Nat Rev Immunol.* 2016;16:626-38, <http://dx.doi.org/10.1038/nri.2016.90>.
97. Malaviya AN, Aggarwal VK, Rawat R, Baghel S, Thakran R, Zaheer Q, et al. Screening for latent tuberculosis infection among patients with rheumatoid arthritis in the era of biologics and targeted synthetic disease-modifying anti-rheumatic drugs in India, a high-burden TB country: The importance of Mantoux and QuantiFERON-TB Gold tests. *Int J Rheum Dis.* 2018;21:1563-71, <http://dx.doi.org/10.1111/1756-185X.13261>.