The relationship between systemic immune inflammation index and disease activity in ankylosing spondylitis patients

Ankilozan spondilit hastalarında sistemik immün inflamasyon indeksinin hastalık aktivitesi ile ilişkisi

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Abstract

Purpose: This study aimed to investigate the relationship between the systemic immune inflammation index (SII), platelet-to-lymphocyte ratio (PLR), neutrophil-to-lymphocyte ratio (NLR), and disease activity and functional status in patients with ankylosing spondylitis (AS).

Materials and methods: This cross-sectional clinical study included a total of 90 patients diagnosed with AS according to the Modified New York Criteria, aged between 18 and 65, who presented to our outpatient clinics. Demographic data and laboratory parameters, including platelet, neutrophil, basophil, eosinophil, and lymphocyte counts, mean platelet volume (MPV), red blood cell distribution width (RDW), C-reactive protein (CRP), and erythrocyte sedimentation rate (ESR), were recorded. NLR and PLR values were calculated. The SII was calculated by dividing the product of neutrophil and platelet counts by the lymphocyte count.

Results: The study included 90 AS patients (mean age: 42.9 ± 11.3 years). Positive correlations were observed between SII and CRP (p=0.010, r=0.269) and ESR (p=0.007, r=0.282). No significant correlations were found between SII and BASDAI (p=0.323), BASFI (p=0.124) or BASMI (p=0.673). NLR and PLR values didn't differ significantly between active and inactive disease groups across all disease activity measures (BASDAI, ASDASCRP, and ASDAS-ESR; NLR: p=0.933, p=0.639, p=0.240; PLR: p=0.708, p=0.858, p=0.351; respectively). There was a significant correlation between SII and Ankylosing Spondylitis Disease Activity Score with erythrocyte sedimentation rate (ASDAS-ESR) (rho=0.282, p=0.007).

Conclusion: The study suggests that SII correlates positively with CRP and ESR, common inflammatory markers in AS. SII could be a potential marker for assessing inflammation, especially in patients with higher disease activity.

Keywords: Ankylosing spondylitis, complete blood count, inflammation.

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Öz

Amaç: Bu çalışma, ankilozan spondilit (AS) hastalarında sistemik immün inflamasyon indeksi (SII), trombositlenfosit oranı (PLR), nötrofil-lenfosit oranı (NLR) ile hastalık aktivitesi ve fonksiyonel durum arasındaki ilişkiyi araştırmayı amaçlamaktadır.

Gereç ve yöntem: Bu klinik kesitsel çalışmaya, Modifiye New York Kriterleri'ne göre AS tanısı almış 18-65 yaş aralığında 90 hasta dahil edildi. Demografik veriler, trombosit, nötrofil, bazofil, eozinofil ve lenfosit sayıları, ortalama trombosit hacmi (MPV), eritrosit dağılım genişliği (RDW), C-reaktif protein (CRP) ve eritrosit sedimantasyon hızı (ESR) gibi laboratuvar parametreleri kaydedildi. NLR, PLR ve SII değerleri hesaplandı. **Bulgular:** Çalışmaya dahil edilen AS hastalarının yaş ortalamaları 42,9±11,3 yıl idi. SII ile CRP (*p*=0,010, r=0,269) ve ESR (*p*=0,007, r=0,282) arasında pozitif korelasyon saptandı. SII ile BASDAI (*p*=0,323), BASFI (*p*=0,124) veya BASMI (*p*=0,673) arasında anlamlı bir ilişki bulunmadı. NLR ve PLR değerleri aktif ve inaktif hastalık grupları arasında anlamlı bir fark göstermedi (BASDAI, ASDAS-CRP ve ASDAS-ESR için sırasıyla; NLR: *p*=0,933, *p*=0,639, *p*=0,240; PLR: *p*=0,708, *p*=0,858, *p*=0,351). SII ile ESR kullanılarak hesaplanan Ankilozan Spondilit Hastalık Aktivite Skoru (ASDAS-ESR) arasında anlamlı bir korelasyon vardı (rho=0,282, *p*=0,007). **Sonuç:** Çalışma, SII'nin AS'de yaygın inflamatuar belirteçler olan CRP ve ESR ile pozitif korelasyon gösterdiğini ortaya koymaktadır. SII, özellikle daha yüksek hastalık aktivitesine sahip hastalarda inflamasyonu değerlendirmek için potansiyel bir belirteç olabilir.

Anahtar kelimeler: Ankilozan spondilit, tam kan sayımı, inflamasyon.

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Introduction

Ankylosing spondylitis (AS) is a chronic inflammatory rheumatic disease belonging to the spondyloarthritis group, primarily affecting the spine and sacroiliac joints [1]. Due to its progressive nature, disease activity needs to be regularly monitored, and the treatment plan should be adjusted according to changes in activity. C-reactive protein (CRP) and erythrocyte sedimentation rate (ESR) are commonly utilized markers for inflammation assessment during follow-ups. However, their elevation can occur due to various non-specific factors like infections, malignancies, and inflammation unrelated to AS. This highlights the necessity for more specific markers in evaluating disease activity in rheumatic diseases like AS and rheumatoid arthritis [2].

Complete blood count is an easily assessable, cost-effective, and straightforward test. There are changes in blood parameters during inflammatory processes, and these changes can be utilized to assess the level of inflammation [2]. The value obtained by dividing the number of neutrophils by the number of lymphocytes in a complete blood count, known as Neutrophil-to-Lymphocyte Ratio (NLR), has been found to be associated with the level of inflammation in diseases such as thyroid disorders [3], inflammatory bowel disease [4], and diabetes mellitus [5]. Another value obtained by dividing the platelet count by the lymphocyte count, known as Platelet-to-Lymphocyte Ratio (PLR), has found applications in conditions such as liver fibrosis [6] and diabetes [7]. The Systemic Immune Inflammation Index (SII), a novel marker, is calculated by multiplying the platelet count by the neutrophil count and then dividing this product by the lymphocyte count. SII has been found to be more successful in determining inflammation compared to NLR and PLR [8]. In studies, SII has been reported as a prognostic and activity determinant in conditions such as various types of malignancies [9, 10], Behçet's disease [11], vasculitis [12], lateral epicondylitis [13], rheumatoid arthritis [14] and post-stroke depression [15].

Recent studies have reported that data obtained through calculations of laboratory parameters such as the SII, NLR, and PLR can potentially be used to assess disease activity in AS, although this remains a topic of debate [16, 17]. Additionally, in the follow-up of AS patients, parameters that include assessments of symptom severity such as Bath Ankylosing Spondylitis Disease Activity Index (BASDAI), the Ankylosing Spondylitis Disease Activity Score with C-reactive protein (ASDAS-CRP), and the Ankylosing Spondylitis Disease Activity Score with erythrocyte sedimentation rate (ASDAS-ESR) hold significant importance. The aim of this study is to evaluate the relationship between SII, PLR, NLR values and BASDAI, ASDAS-CRP, and ASDAS-ESR scores, which are used to determine disease activity and their applicability in patient monitoring among AS patients.

Materials and methods

Our study was designed as a cross-sectional clinical study. Between May 2, 2024, and May 20, 2024, a total of 90 patients between the ages of 18-65 who were diagnosed with AS according to the Modified New York Criteria and followed up at the Health Science University, Istanbul Physical Therapy and Rehabilitation Training and Research Hospital and Beylikdüzü State Hospital outpatient clinics were included in the study. Ethical clearance for this research, as per protocol number 2024/18, was granted by the Clinical Research Ethics Committee of Istanbul Physical Medicine and Rehabilitation Training and Research Hospital on April 30, 2024. Prior to the commencement of the study, participants provided informed consent. The study was conducted in accordance with the guidelines outlined in the Declaration of Helsinki.

Exclusion criteria for our study included the presence of acute or chronic infections, autoimmune diseases other than ankylosing spondylitis, pregnancy, diabetes, chronic kidney or liver disease, and the presence of cardiovascular or hematological diseases that could lead to changes in laboratory parameters. At the beginning of the study, sociodemographic data such as age, gender, marital status, bodymass index (BMI), smoking status, duration of disease diagnosis, and age at symptom onset were recorded for all patients.

Disease activity was assessed using the BASDAI, ASDAS-CRP, and ASDAS-ESR. The Turkish version of BASDAI, which has been validated and demonstrated reliability by Akkoç et al. [18], was utilized in the study. Scores of 4 or higher were considered indicative of active disease, while scores below 4 were classified as inactive disease. For ASDAS-CRP and ASDAS-ESR scores, patients with values of 2.1 and above were categorized as active, while those with values below 2.1 were placed in the inactive group [19]. Spinal mobility was evaluated using the Bath Ankylosing Spondylitis Metrology Index (BASMI), and functional status was assessed using the Bath Ankylosing Spondylitis Functional Index (BASFI) [20]. Furthermore, during the routine follow-up of patients, requested blood tests were examined record platelet, neutrophil, basophil, eosinophil, lymphocyte counts, mean platelet volume (MPV), red blood distribution width (RDW), CRP, and ESR values. NLR, PLR and SII were calculated. SII is obtained by dividing the product of neutrophil and platelet counts by the lymphocyte count.

Statistical analysis

The study's sample size was determined based on Wu et al. [21] research, with correlations of 0.483 for SII and CRP, 0.374 for SII and ESR, and 0.667 for SII and BASDAI. With a significance level of 5% and 95% power, sample sizes for CRP, ESR, and BASDAI were calculated as 41, 73, and 19, respectively. A minimum of 73 participants was needed for all variables. Our study included 90 AS patients.

We used G*Power 3.1.9.4 for sample size calculation. Data normality was assessed with Kolmogorov-Smirnov test. Descriptive statistics were used for quantitative data, presented as mean/standard deviation or median/min-max, and categorical data as frequency/percentage. Spearman test analyzed SII-AS disease activity correlation due to non-normality. Mann-Whitney U test compared active and inactive disease groups. SPSS 21.0 conducted statistical analyses.

Results

Our study included 90 patients diagnosed with AS. Detailed descriptive statistics are provided in Table 1. Of the participants, 61.1% were male, and 38.9% were female, with a median age of 45.5. The median BMI of the individuals was 27.7, with a minimum value of 11.5 and a maximum value of 41.6 (Table 1).

Table 2 illustrates the correlation between AS disease activity parameters and laboratory data. SII doesn't show significant relationships with BASDAI (p=0.323), BASFI (p=0.124), or BASMI scores (p=0.673), but it correlates positively with CRP and ESR values (p=0.010, r=0.269 and p=0.007, r=0.282, respectively). PLR and NLR ratios aren't correlated with any parameter. BASMI scores correlate significantly with CRP (p=0.026) and ESR (p=0.005), whereas BASFI and BASDAI scores don't exhibit such correlations. As expected, CRP and ESR values correlate with ASDAS-CRP (p=0.000 and p=0.000, respectively) and ASDAS-ESR scores (p=0.014 and p=0.000, respectively) (Table 2).

In Table 3, patients were examined in two groups based on BASDAI, ASDAS-CRP, and ASDAS-ESR scores, classified as active and inactive. The calculated SII value in patients with high ASDAS-ESR scores, indicating active disease, was significantly higher than in the inactive group (p=0.032). There was no significant difference in NLR (p=0.933, p=0.639, p=0.240) and PLR (p=0.708, p=0.858, p=0.351) between the groups across all disease activity measures (BASDAI, ASDAS-CRP, and ASDAS-ESR, respectively) (Table 3).

Table 1. Descriptive statistics

		Median (min-max) / n (%)	%
Age		45.5 (21-65)	
Gender	Male	55	61.1
Gender	Female	35	38.9
Marital status	Married	68	75.6
Waritai Status	Single	22	24.4
ВМІ		27.7 (11.5-41.6)	
	Non-smoker	48	53.3
C making	<10 pack-years	13	14.4
Smoking	10-20 pack-years	15	16.7
	>20 pack-years	14	15.6
Duration of diagnosis		84 (1-396)	
Age at symptom onset		24.5 (8-57)	
LI A DOZ	Positive	51	56.7
HLA B27	Negative	39	43.3
BASDAI		5 (0-10)	
BASFI		4 (0-10)	
BASMI		2 (0-8)	
CRP		3.9 (0.2-46.4)	
ESR		9 (2-60)	
MPV		9.2 (7.1-12.5)	
RDW		13.4 (11.9-21.2)	
NLR		1.7 (0.8-4.6)	
PLR		111 (45.6-258)	
SII		478 (198.6-1503.7)	
ASDAS-CRP		2.7 (1.1-5.2)	
ASDAS-ESR		2.7 (1-5)	

BMI: Body-mass index BASDAI: Bath Ankylosing Spondylitis Disease Activity Index, BASFI: Bath Ankylosing Spondylitis Functional Index BASMI: Bath Ankylosing Spondylitis Metrology Index, ASDAS-CRP: Ankylosing Spondylitis Disease Activity Score with C-reactive protein ASDAS-ESR: Ankylosing Spondylitis Disease Activity Score with erythrocyte sedimentation rate, CRP: C-reactive protein ESR: Erythrocyte sedimentation rate, PLR: platelet to lymphocyte ratio, NLR: neutrophil to lymphocyte ratio SII: systemic immune inflammation index

Table 2. Correlation between laboratory parameters and parameters related to disease activity

	ВА	BASDAI	BA	BASMI	ВА	BASFI	S	CRP	ш	ESR	ASDA	ASDAS-CRP	ASD,	ASDAS-ESR
	rho	р	rho	d	rho	d	rho	d	rho	d	rho	d	rho	d
CRP	-0.029	0.785	0.234	0.026*	0.072	0.499	-	ı	0.696	*000.0	0.473	*000.0	0.258	0.014*
ESR	0.016	0.879	0.296	0.005*	0.010	0.926	969.0	*000.0	-		0.374	*000.0	0.450	*000.0
RDW	-0.061	0.568	0.199	0.060	-0.100	0.348	0.124	0.245	0.315	0.003*	0.055	0.605	0.131	0.218
MPV	-0.049	0.649	0.041	0.701	-0.111	0.296	0.002	0.986	0.001	0.990	-0.009	0.932	-0.021	0.845
S	0.105	0.323	-0.045	0.673	0.163	0.124	0.269	0.010*	0.282	0.007*	0.196	0.064	0.196	0.065
PLR	0.103	0.335	-0.019	0.860	0.112	0.295	-0.006	0.953	0.133	0.212	0.031	0.771	0.098	0.357
NLR	-0.007	0.949	-0.081	0.446	0.097	0.361	0.146	0.171	0.109	0.308	0.031	0.770	0.007	0.948

Spearman correlation test. Values with p<0.05 are marked with an asterisk (*). BASDAI: Bath Ankylosing Spondylitis Disease Activity Index, BASFI: Bath Ankylosing Spondylitis Functional Index BASMI: Bath Ankylosing Spondylitis Metrology Index, CRP: C-reactive protein, ASDAS-CRP: Ankylosing Spondylitis Disease Activity Score with erythrocyte sedimentation rate, ESR: Erythrocyte sedimentation rate, PLR: platelet to lymphocyte ratio NLR: neutrophil to lymphocyte ratio, SII: systemic immune inflammation index

Table 3. Changes in laboratory and clinical parameters according to disease activity

		BASDAI				ASDAS-CRP			AS	ASDAS-ESR		
	<4 (n=27)	≥4 (n=63)	۵	z	<2.1 (n=22)	≥2.1 (n=68)	Q	z	<2.1 (n=25)	≥2.1 (n=65)	d	×
BASFI	BASFI 1 (0-6)	5 (1-10)	*000.0	0.000* -5.427	1.5 (0-6)	5 (0-10)	*000.0	0.000* -3.524 1 (0-7)	1 (0-7)	5 (0-10)	*000.0	-3.912
BASMI	3 (0-6)	2 (0-8)	0.535	-0.621	2 (0-6)	2 (0-8)	0.404	-0.835	2 (0-6)	2 (0-8)	0.967	-0.041
RDW	13.4 (12.5-19.6)	13.4 (11.9-21.2)	0.558	-0.586	13.2 (12.5-19.6)	13.4 (11.9-21.2)	0.305	-1.025	-1.025 13.4 (12.5-19.6)	13.4 (11.9-21.2)	0.708	-0.374
MPV	9.2 (7.2-10.7)	9.1 (7.1-12.5)	0.754	-0.313	9.2 (8-10.1)	9.1 (7.1-12.5)	0.764	-0.301	-0.301 9.2 (7.8-10.1)	9.2 (7.1-12.5)	0.435	-0.780
CRP	4.2 (0.2-36.6)	3.6 (0.2-46.4)	0.853	-0.185	1.8 (0.2-12.2)	4.9 (0.2-46.4)	0.002*	-3.108	2.4 (0.3-23.1)	4.7 (0.2-46.4)	0.051	-1.951
ESR	8 (2-51)	10 (2-60)	0.363	-0.910	7 (2-35)	11 (2-60)	0.042*	-2.035	-2.035 6 (2-35)	11 (2-60)	*200.0	-2.703
NLR	1.6 (0.8-4.4)	1.8 (0.8-4.6)	0.933	-0.084	1.6 (0.8-4.4)	1.76 (0.8-4.6)	0.639	-0.469	1.6 (0.9-4.4)	1.8 (0.8-4.6)	0.240	-1.176
PLR	105.7 (60.9-223.6)	111.6 (45.6-258)	0.708	-0.374	110.1 (60.9-195.3)	110.9 (45.6-258)	0.858	-0.178	105.7 (60.9-165.2) 111.8 (45.6-257.9)	111.8 (45.6-257.9)	0.351	-0.932
S	438.7 (214.6-1017.4)	438.7 (214.6-1017.4) 496 (198.6-1503.7) 0.410	0.410	-0.823	412.8 (205-847.3)	491.12 (198.6-1503.7) 0.233		-1.192	357.4 (205-847.3)	357.4 (205-847.3) 496 (198.6-1503.7) 0.032* -2.139	0.032*	-2.139

Mann Whitney U test. Values with ρ<0.05 are marked with an asterisk (*). Bath Ankylosing Spondylitis Disease Activity Index, BASFI: Bath Ankylosing Spondylitis Functional Index, BASMI: Bath Ankylosing Spondylitis Disease Activity Score with C-reactive protein , ASDAS-ESR: Ankylosing Spondylitis Disease Activity Score with erythrocyte sedimentation rate, CRP: C-reactive protein ESR: Erythrocyte sedimentation rate, PLR: platelet to lymphocyte ratio, NLR: neutrophil to lymphocyte ratio, SII: systemic immune inflammation index

Discussion

In this cross-sectional clinical study, a positive correlation was found between CRP and ESR values, which are frequently used in the follow-up and treatment decisions of AS patients and considered as markers of inflammation, and the SII value. Additionally, higher SII values were found in the patient group considered active according to the ASDAS-ESR score. However, there was no significant difference in SII values between active and inactive patient groups based on ASDAS-CRP and BASDAI scores. Furthermore, in our study, we did not find a significant difference in NLR and PLR levels between active and inactive groups for all three parameters.

There are many studies in the literature investigating the use of blood parameters to determine disease activity and periodic patient monitoring in rheumatological diseases [14, 15]. However, the number of such studies in AS patients is limited. In the follow-up and determination of activity in AS patients, not only laboratory parameters but also scoring systems such as BASDAI, ASDAS-CRP, and ASDAS-ESR are used. Limitation in spinal mobility can be evaluated with the BASMI index, and the patient's functional status can be assessed with the BASFI scoring. Scoring systems that rely on the patient's self-report, such as BASDAI and BASFI, are influenced by many additional factors beyond the inflammation caused by the disease, including the patient's psychological state, perception of the disease, and central sensitization. ASDAS-CRP and ASDAS-ESR values include both laboratory parameters and patient self-report when calculated [22]. The lack of correlation between SII and scoring systems obtained from subjective questioning, but its higher values in patients considered active according to ASDAS-ESR, may be attributed to the patient's mental and psychological factors. The correlation of SII with ESR and CRP values also suggests that it could be a biomarker with the potential to predict the current level of inflammation.

In a study, the SII value in AS patients was found to be correlated with ESR, CRP, and BASDAI scores [21]. In another study, SLE, RA, and AS patient groups were compared with healthy controls. SII values were higher in AS and RA patients compared to the control group.

PLR was higher in all three groups, while NLR was significantly higher only in SLE patients compared to the control group. However, when the AS group was grouped according to the level of disease activity, there was no significant difference in SII, NLR, and PLR values. Additionally, in AS patients, SII and NLR values were correlated with CRP, ESR, and ASDAS parameters, while PLR was not correlated. On the other hand, MPV and RDW values showed changes consistent with disease activity in all three disease groups [23]. In contrast, in our study, RDW and MPV values in AS patients were not found to be associated with disease activity or laboratory parameters.

In the study by Liang et al. [24], PLR and NLR ratios were significantly higher in the AS patient group compared to the healthy group. Additionally, the AS patient group was categorized as active and inactive based on BASDAI scores, and the active group had significantly higher PLR values, but the same was not true for NLR. In our research, parallel to this study, NLR values were not found to be associated with disease activity, and similarly, PLR values were not found to be different between active and inactive patients, as was the case with NLR. In a study conducted by Osami et al. [25] in AS patients, NLR, PLR, and ESR values were found to be significantly higher in active AS patients compared to inactive ones. However, when compared to the healthy group, ESR was significantly higher, while NLR and PLR ratios were similar between the two groups. In contrast to this study, in our research, NLR and PLR values were not correlated with disease activity levels indicated by BASDAI, ASDAS-CRP, and ASDAS-ESR values.

Our study was conducted only with AS-diagnosed patients due to the absence of a control group. Additionally, the single-center nature of the study is one of its limitations. One of the strengths of our research is that we examined the correlation of markers by simultaneously using multiple methods for evaluating disease activity. The markers evaluated in this study are cost-effective parameters that can be easily obtained in all laboratories where complete blood counts are performed and are well-correlated with inflammation [23]. In our study, especially, SII has been shown to be a cost-effective marker that can be used for this

purpose. Further studies with a larger number of patients and multi-center studies are needed to establish its safe use in patient follow-up.

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References

- Rudwaleit M, Baeten D. Ankylosing spondylitis and bowel disease. Best Pract Res Clin Rheumatol. 2006;20(3):451-471. doi:10.1016/j.berh.2006.03.010
- Colglazier CL, Sutej PG. Laboratory testing in the rheumatic diseases: a practical review. South Med J. 2005;98(2):185-191. doi:10.1097/01. SMJ.0000153572.22346.E9
- Aktas G, Sit M, Dikbas O, et al. Elevated neutrophil-to-lymphocyte ratio in the diagnosis of Hashimoto's thyroiditis. Rev Assoc Med Bras (1992). 2017;63(12):1065-1068. doi:10.1590/1806-9282.63.12.1065
- Posul E, Yilmaz B, Aktas G, Kurt M. Does neutrophil-tolymphocyte ratio predict active ulcerative colitis?. Wien Klin Wochenschr. 2015;127(7-8):262-265. doi:10.1007/ s00508-014-0683-5
- Duman TT, Aktas G, Atak BM, Kocak MZ, Erkus E, Savli H. Neutrophil to lymphocyte ratio as an indicative of diabetic control level in type 2 diabetes mellitus. *Afr Health Sci.* 2019;19(1):1602-1606. doi:10.4314/ahs. v19i1.35
- Kosekli MA. Mean platelet volume and platelet to lymphocyte count ratio are associated with hepatitis B-related liver fibrosis. Eur J Gastroenterol Hepatol. 2022;34(3):324-327. doi:10.1097/ MEG.00000000000002219
- Atak B, Aktas G, Duman TT, Erkus E, Kocak MZ, Savli H. Diabetes control could through platelet-tolymphocyte ratio in hemograms. Rev Assoc Med Bras (1992). 2019;65(1):38-42. doi:10.1590/1806-9282.65.1.38

- Aziz MH, Sideras K, Aziz NA, et al. The Systemicimmune-inflammation Index Independently Predicts Survival and Recurrence in Resectable Pancreatic Cancer and its Prognostic Value Depends on Bilirubin Levels: A Retrospective Multicenter Cohort Study. Ann Surg. 2019;270(1):139-146. doi:10.1097/ SLA.0000000000000000660
- De Giorgi U, Procopio G, Giannarelli D, et al. Association of Systemic Inflammation Index and Body Mass Index with Survival in Patients with Renal Cell Cancer Treated with Nivolumab. Clin Cancer Res. 2019;25(13):3839-3846. doi:10.1158/1078-0432.CCR-18-3661
- Jomrich G, Paireder M, Kristo I, et al. High Systemic Immune-Inflammation Index is an Adverse Prognostic Factor for Patients With Gastroesophageal Adenocarcinoma. *Ann Surg.* 2021;273(3):532-541. doi:10.1097/SLA.000000000003370
- Tanacan E, Dincer D, Erdogan FG, Gurler A. A cutoff value for the Systemic Immune-Inflammation Index in determining activity of Behçet disease. Clin Exp Dermatol. 2021;46(2):286-291. doi:10.1111/ced.14432
- Kim Y, Choi H, Jung SM, Song JJ, Park YB, Lee SW. Systemic immune-inflammation index could estimate the cross-sectional high activity and the poor outcomes in immunosuppressive drug-naïve patients with antineutrophil cytoplasmic antibody-associated vasculitis. Nephrology (Carlton). 2019;24(7):711-717. doi:10.1111/nep.13491
- Uysal A. Investigating the presence of inflammation in lateral epicondylitis with platelet/lymphocyte ratio, neutrophil/lymphocyte ratio, and systemic immuneinflammation index. *Chron Precis Med Res.* 2023;4:29-33. doi:10.5281/zenodo.7718532
- Choe JY, Lee CU, Kim SK. Association between Novel Hematological Indices and Measures of Disease Activity in Patients with Rheumatoid Arthritis. *Medicina* (*Kaunas*). 2023;59(1):117. Published 2023 Jan 6. doi:10.3390/medicina59010117
- Satis S. New Inflammatory Marker Associated with Disease Activity in Rheumatoid Arthritis: The Systemic Immune-Inflammation Index. Curr Health Sci J. 2021;47(4):553-557. doi:10.12865/CHSJ.47.04.11
- Song GG, Lee YH. Red cell distribution width, platelet-to-lymphocyte ratio, and mean platelet volume in ankylosing spondylitis and their correlations with inflammation: A meta-analysis. Mod Rheumatol. 2020;30(5):894-899. doi:10.1080/14397595.2019.164 5373
- Khorrampazhouh N, Omranzadeh A, Fazeli B, et al. A Systematic Review and Meta-analysis of Clinical Studies on Ankylosing Spondylitis and Neutrophil to Lymphocyte Ratio. *Curr Rheumatol Rev.* 2022;18(2):160-167. doi:10.2174/1573397117666210 921114431

- Akkoc Y, Karatepe AG, Akar S, Kirazli Y, Akkoc N. A
 Turkish version of the Bath Ankylosing Spondylitis
 Disease Activity Index: reliability and validity. Rheumatol
 Int. 2005;25(4):280-284. doi:10.1007/s00296-0030432-y
- Machado PM, Landewé R, Heijde DV; Assessment of SpondyloArthritis international Society (ASAS). Ankylosing Spondylitis Disease Activity Score (ASDAS): 2018 update of the nomenclature for disease activity states. *Ann Rheum Dis*. 2018;77(10):1539-1540. doi:10.1136/annrheumdis-2018-213184
- Ozer HT, Sarpel T, Gulek B, Alparslan ZN, Erken E. The Turkish version of the Bath Ankylosing Spondylitis Functional Index: reliability and validity. *Clin Rheumatol*. 2005;24(2):123-128. doi:10.1007/s10067-004-0984-6
- Wu J, Yan L, Chai K. Systemic immune-inflammation index is associated with disease activity in patients with ankylosing spondylitis. *J Clin Lab Anal*. 2021;35(9):e23964. doi:10.1002/jcla.23964
- Sariyildiz A, Benlidayi IC, Turk I, Acemoglu SSZ, Unal I. Evaluation of the relationship between blood cell markers and inflammation, disease activity, and general health status in ankylosing spondylitis. Rev Assoc Med Bras (1992). 2023;69(10):e20230722. Published 2023 Sep 18. doi:10.1590/1806-9282.20230722
- Taha SI, Samaan SF, Ibrahim RA, Moustafa NM, El Sehsah EM, Youssef MK. Can Complete Blood Count Picture Tell Us More About the Activity of Rheumatological Diseases?. Clin Med Insights Arthritis Musculoskelet Disord. 2022;15:11795441221089182. Published 2022 Apr 22. doi:10.1177/11795441221089182
- 24. Liang T, Chen J, Xu G, et al. Platelet-to-Lymphocyte Ratio as an Independent Factor Was Associated With the Severity of Ankylosing Spondylitis. Front Immunol. 2021;12:760214. Published 2021 Nov 5. doi:10.3389/ fimmu.2021.760214
- Al Osami MH, Awadh NI, Khalid KB, Awadh AI. Neutrophil/lymphocyte and platelet/lymphocyte ratios as potential markers of disease activity in patients with Ankylosing spondylitis: a case-control study. Adv Rheumatol. 2020;60(1):13. Published 2020 Jan 29. doi:10.1186/s42358-020-0113-5