



Analysis of Personality Trait in Patients with Alopecia Areata

Jeong-Min Kim, Hoon-Soo Kim^{1,2}, Hyun-Chang Ko, Byung-Soo Kim^{1,2}, Moon-Bum Kim^{1,2}

Department of Dermatology, Pusan National University Yangsan Hospital, Yangsan, ¹Department of Dermatology, Pusan National University School of Medicine, Busan, ²Bio-Medical Research Institute, Pusan National University Hospital, Busan, Korea

Dear Editor:

Alopecia areata (AA) can be affected by various etiologic factors including immunological and endocrine abnormalities, as well as genetic factors or psychiatric disturbances. In the context of psychiatric disturbances in patients with AA, there have only been a few studies focusing on personality traits. We guess most dermatologists have prejudice about personality trait that AA patients may have negative personality traits such as antisocial, hysterical, and anxious ones, though they can be primary or secondary. Güleç et al.¹ reported that certain personality traits can modulate the onset of AA. Additionally, Carrizosa et al.² used Minnesota Multiphasic Personality Inventory to show that AA patients express more depressive, hysterical and anxious feelings than healthy subjects. However, Alfani et al.³ recently reported that patients with AA do not have specific personality traits, as shown through an assessment using the International Personality Disorder Examination. Considering these findings, the personality traits associated with AA seems to be equivocal. Therefore, we aimed to clarify whether there are specific personality traits associated with AA patients to evaluate the correlation between prognostic factors of AA and personality traits under the assumption that these patients have a specific personality traits and that negative prognostic factors of AA can affect the formation of more negative personality traits than positive ones. One hundred patients with AA and 100 healthy subjects (control group) were included in this study. Between the two groups, there were no statistical differences in terms of sex, age or ethnicity

that might reasonably be expected to have an influence on the onset of AA. After age- and sex-matching the subjects from the two groups, both patients with AA and healthy controls had consultations at our dermatologic clinic. Before the start of the study, all subjects were screened for comorbid psychiatric conditions and none of them had ever had a psychiatric diagnosis⁴. All subjects were asked to fill-out a NEO-Five Factor Inventor (NEO-FFI) questionnaire, the most commonly used psychological personality inventory and was developed and revised by De Fruyt et al.⁵. This inventory can be used to assess five personality domains of neuroticism (N), extraversion (E), openness (O), agreeableness (A) and conscientiousness (C) through a more convenient and valid method compared to that used in the original five-factor approach. All statistical analyses were performed through 2-sample t-tests using PASW Statistics ver. 18.0 software (IBM Co., Armonk, NY, USA). Comparison were made not only between the scores of the AA and control groups but also between those of 2 AA groups based on AA prognostic factors; onset age around adolescence (≤ 13 vs. > 13 years)⁶⁻⁸, duration (≤ 2 vs. > 2 years)⁹, severity ($\leq 50\%$ vs. $> 50\%$ involvement of scalp)¹⁰, and involvement of nails⁷. Contrary to our expectations, the E ($p=0.013$) and A ($p<0.001$) domains of the NEO-FFI were more dominant in the AA group than in controls. Furthermore, no significant difference were found between the prognostic factors of AA and personality traits, except for a significantly lower trait A value for patients whose disease duration was over 2 years. Overall, there was no significant difference be-

Received August 16, 2016, Revised October 28, 2016, Accepted for publication November 16, 2016

Corresponding author: Moon-Bum Kim, Department of Dermatology, Pusan National University School of Medicine, 305 Gudeok-ro, Seo-gu, Busan 49234, Korea. Tel: 82-51-240-7338, Fax: 82-51-245-9467, E-mail: drkmp@hanmail.net

This is an Open Access article distributed under the terms of the Creative Commons Attribution Non-Commercial License (<http://creativecommons.org/licenses/by-nc/4.0>) which permits unrestricted non-commercial use, distribution, and reproduction in any medium, provided the original work is properly cited.

Copyright © The Korean Dermatological Association and The Korean Society for Investigative Dermatology

Table 1. Average NEO-FFI scores in good and poor prognostic groups in relation to each prognostic factor

Prognosis factor	Domains	Good prognostic group	Poor prognostic group	<i>p</i> -value
Onset age [†]	N	35.43	36.18	0.678
	E	43.00	38.32	0.344
	O	36.14	36.83	0.849
	A	24.86	26.98	0.112
	C	37.71	38.48	0.336
Disease duration [‡]	N	36.03	36.75	0.934
	E	38.50	39.67	0.922
	O	36.84	36.67	0.201
	A	36.97	26.02	0.041*
Disease severity [§]	C	38.29	39.42	0.464
	N	35.95	36.73	0.944
	E	38.64	38.64	0.188
	O	36.79	36.91	0.769
Nail change	A	26.81	27.09	0.426
	C	38.97	36.50	0.093
	N	34.14	36.26	0.372
	E	38.43	38.66	0.991
	O	38.28	36.71	0.141
	A	34.00	27.08	0.148
	C	40.71	38.26	0.319

Good prognostic group vs. poor prognostic group: [†] ≤13 (n=7) vs. >13 years (n=93), [‡] ≤2 (n=89) vs. >2 years (n=11), [§] ≤50% (n=79) vs. >50% (n=21), ^{||} with nail change (n=93) vs. without nail change (n=7). NEO-FFI: NEO-Five Factor Inventor, N: neuroticism, E: extraversion, O: openness, A: agreeableness, C: conscientiousness. **p*-values less than 0.05 were considered significant.

tween the personality of the AA and control groups, as a whole (Table 1). Although the questionnaires and the populations from which patients and controls were selected for this study differed from those used in previous studies, we believe that our results are meaningful because they are supported by 2 distinct comparisons, namely, between AA and control groups and between 2 AA groups with different AA prognostic factors. This is the first trial that has attempted to define the relationship between personality traits associated with AA and prognostic factors of AA. Considering our findings, we conclude that negative personality traits such as such as antisocial behavior, hysteria, and anxiety are not characteristic features of patients with AA. Therefore, we agree with Alfani et al.³ in believing

that patients with AA do not have specific personality traits. Thus, we need to get rid of prejudice that patients with AA have negative or specific personality traits during treatment from present study.

CONFLICTS OF INTEREST

The authors have nothing to disclose.

REFERENCES

- Güleç AT, Tanriverdi N, Dürü C, Saray Y, Akçali C. The role of psychological factors in alopecia areata and the impact of the disease on the quality of life. *Int J Dermatol* 2004;43:352-356.
- Carrizosa A, Estepa-Zabala B, Fernandez-Abascal B, Garcia-Hernandez MJ, Ruiz-Dablado S. Alopecia areata: a specific personality? *Int J Dermatol* 2005;44:437-438.
- Alfani S, Antinone V, Mozzetta A, Di Pietro C, Mazzanti C, Stella P, et al. Psychological status of patients with alopecia areata. *Acta Derm Venereol* 2012;92:304-306.
- Lee NR, Kim BK, Yoon NY, Lee SY, Ahn SY, Lee WS. Differences in comorbidity profiles between early-onset and late-onset alopecia areata patients: a retrospective study of 871 Korean patients. *Ann Dermatol* 2014;26:722-726.
- De Fruyt F, McCrae RR, Szirmák Z, Nagy J. The five-factor personality inventory as a measure of the five-factor model: Belgian, American, and Hungarian comparisons with the NEO-PI-R. *Assessment* 2004;11:207-215.
- You HR, Kim SJ. Factors associated with severity of alopecia areata. *Ann Dermatol* 2017;29:565-570.
- Otberg N, Shapiro J. Hair growth disorders. In: Goldsmith LA, Kats SI, Gilchrist BA, Paller AS, Leffell DJ, Wolff K, editors. *Fitzpatrick's dermatology in general medicine*. 8th ed. New York: McGraw-Hill, 2012:991-995.
- Cho HH, Jo SJ, Paik SH, Jeon HC, Kim KH, Eun HC, et al. Clinical characteristics and prognostic factors in early-onset alopecia totalis and alopecia universalis. *J Korean Med Sci* 2012;27:799-802.
- Olsen EA. Investigative guidelines for alopecia areata. *Dermatol Ther* 2011;24:311-319.
- Tosti A, Bellavista S, Iorizzo M. Alopecia areata: a long term follow-up study of 191 patients. *J Am Acad Dermatol* 2006;55:438-441.