

Study of Skin Expression of MicroRNA-146a in Patients with Alopecia Areata

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Abstract

Alopecia areata [AA] may be an immune system issue portrayed Toward transient, non-scarring hair reduction What's more protection of the hair follicle. Hair reduction might detract numerous structures going from reduction in well-defined patches with diffuse alternately aggregate hair loss, which might influence every last bit hair bearing locales. MicroRNAs [miRNAs] are little noncoding rna atoms ~20 nucleotides long. Furthermore they need developed Concerning illustration critical translational controllers for genes over Different tissues Furthermore living procedures included clinched alongside immune system reactions. MicroRNA-146a may be found in the LOC285628 gene ahead mankind's chromosome 5. It will be communicated predominantly done resistant tissues, Also its outflow camwood be prompted upon resistant Mobile development or actuation. Point of the work: is with explore the part of miRNA 146a done pathogenesis from claiming AA Subjects Also .four mm punch biopsies were made from each subject. MiRNA needed been concentrated What's more miR-146a needed been assessed altogether tests utilizing RT-qPCR. Every one techniques were performed as stated by the manufacturer's guidelines. there might have been statistically noteworthy distinction between patients one assembly What's more control aggregations in regards to miRNA 146a outflow level. Conclusion: miRNA-146a is fundamentally upregulated in AA which helps the idea from claiming epigenetic inclusion On AA pathogenesis.

Keywords: Alopecia areata, MiRNA-146a.

1. Introduction

Alopecia areata [AA] may be a immune system issue described by transient, non-scarring hair passing Also protection of the hair follicle. Hair passing might detract numerous structures going from reduction Previously, well-defined patches with diffuse or downright hair loss, which might influence every one hair bearing destinations. AA influences almost 2% of the all populace Sooner or later Throughout their life the long haul [17].

Same time those correct etiopathogenesis of AA stays unclear, there is respectable confirmation suggesting immune system [12], hereditary [11] to ailment pathogenesis.

Differential levels about circle miRNAs bring been identifier to immune system ailments imparting comparative pathogenic pathways will AA, for example, such that rheumatoid joint inflammation [3] , vitiligo Also psoriasis[6]. Therefore, miRNAs might serve possibility parts as symptomatic markers and workable restorative focuses done AA and also blacks [23].

MicroRNA-146a will be found in the LOC285628 gene ahead human chromosome 5, responds with lipopolysaccharide [LPS] incitement Previously, mankind's monocytes and the incitement for miR-146a will be atomic component κ B [NF- κ B] reliant [22].

Investigations bring shown that miR-146a assumes pivotal parts in the pathogenesis for immune system diseases, for example, such that systemic lupus erythematosus [SLE] [24], rheumatoid joint inflammation [RA] [13], and Sjögren's syndrome [SS] [15]. However, its part for AA may be not broadly investigated.

2. Subjects and Methods

This case control study was conducted on thirty patients [group A] with different clinical variants of AA. In addition, twenty apparently healthy individuals of matched age and sex were chosen as a control group [group B]. Patients were recruited from outpatient clinic of Dermatology, Venereology and Andrology Department of Benha University Hospitals.

About 4 mm punch biopsies were taken from each subject and immediately put it in liquid nitrogen and stored at -80°C until tested . MicroRNA was extracted and miRNA-146a was evaluated in all samples using RT-qPCR.

2.1. Statistical analyses

Those gathered information might have been revised, coded, tabulated Furthermore presented to a pc utilizing Factual one bundle to social science [IBM corp. Discharged 2011. IBM SPSS detail for Windows, form 20. 0. Armonk, NY: IBM corp.]. Information were exhibited Furthermore suitability examination might have been completed as stated by those kind of information gotten to every parameter. Unmitigated information were exhibited Similarly as number Furthermore rates same time quantitative .

information were communicated Similarly as intend \pm standard deviation [SD], and range.

3. Results

Table [1] Shows patients with AA had significantly higher miRNA-146a expression level when compared to control group [P<0.05].

4. Discussion

Alopecia areata [AA] will be An nonscarring hair misfortune jumble with a capricious course Furthermore An totally range for manifestations [4].

The miRNA 146a aberrant statement is connected with Different incendiary issue for example, RA What's more psoriasis. An muscle to about

confirmation uncovered upregulation from claiming miRNA 146a done immune system illnesses including AA; the helter skelter statement from claiming miRNA 146a helped immune system infections pathogenesis Eventually Tom's perusing over-activation of the IFN pathway [5].

Table (1) Comparison between patients and control regarding miRNA-146a expression level.

Micro RNA-146a expression level [FC]	Mean± SD	ContrN=20		AAN=30		<0.05
		2.12	±0.023	6.2	±75	
		0.263		3.4		
	Median					

Comes about of the display consider demonstrated huge upregulation of miRNA 146a statement level [4. 3 FC] done scalp biopsy from claiming AA patients in examination will control group, Furthermore also there might have been An sure huge correspondence between its level and the seriousness of the sickness communicated in the manifestation of salt score.

The effects of the available investigation might a chance to be clarified Toward T-cell co-stimulatory signs dysregulation. MiRNA-induced dysregulation for ICOS [Inducible T-cell CO-Stimulator] need been identifier to RA On people [20]. To AA for mice, dependent upon regulation of miRNA 146a predicted to target ICOS. Notably, it might have been distinguished relating huge upregulation of ICOS gene interpretation exhibit. This Might propose that higher statement about miRNA 146a clinched alongside AA skin brought about constitutive outflow for ICOS, which prompts improved T-cell reactions [23].

5. Conclusion

miRNA-146a may be essentially upregulated done AA which backs the idea from claiming epigenetic association clinched alongside AA pathogenesis.

Reference

- [1] G.Ayeldeen, Y.Nassar, H.Ahmed, Possible use of miRNAs-146a and-499 expression and their polymorphisms as diagnostic markers for rheumatoid arthritis, *Mol Cell Biochem*, Vol.449,PP.145-156, 2018.
- [2] O.Bilgic, A.Sivrikaya, A.Unlu , Serum cytokine and chemokine profiles in patients with alopecia areata, *J Dermatolog Treat*,Vol. 27(3),PP. 260-263,2016.
- [3] S.Bluml, M.Bonelli, B.Niederreiter, Essential role of microRNA-155 in the pathogenesis of autoimmune arthritis in mice,*Arthritis Rheum*,Vol. 63,PP.1281–1288, 2011.
- [4] E.Darwin, P.A.Hirt, R.Fertig,B .Doliner, Alopecia areata, Review of epidemiology, clinical features, pathogenesis, and new treatment options, *Int J Trichology*,Vol.10,PP. 51-60, 2018.

- [5] Q.Duan, X.Mao, Y.Xiao, Super enhancers at the miR-146a and miR-155 genes contribute to self-regulation of inflammation, *Biochim Biophys Acta*,Vol.1859,PP. 564–571.2016.
- [6] A.C.V.Fricke and M.Miteva , Epidemiology and burden of alopecia areata, a systematic review, *Clin Cosmet Investig Dermatol*,Vol.8, PP. 397–403, 2015.
- [7] E.Kasumagic Halilovic, A.Prohic and S.Cavaljuga, Tumor necrosis factor-alpha in patients with alopecia areata, *Indian J Dermatol*, Vol.56,PP.494-502, 2011.
- [8] M.B.Lovendorf , H.Mitsui, J.R.Zibert , Laser capture micro dissection followed by next-generation sequencing identifies disease-related micro RNA s in psoriatic skin that reflects systemic micro RNA changes in psoriasis, *Exp Dermatol*,Vol.24,PP. 187-193, 2015.
- [9] L.F.Lu, M.P.Boldin, A. Chaudhry, K.D.Taganov , Function of miR-146a in controlling Treg cell-mediated regulation of Th1 responses, *Cell*,Vol.142,PP.914-929, 2010.
- [10] A.D.McClelland and P.Kantharidis, MicroRNA in the development of diabetic complications, *Clin Sci*, Vol.126,PP.95-110, 2014.
- [11] A.J.McDonagh and R.Tazi-Ahnini, Epidemiology and genetics of alopecia areata, *Clin Exp Dermatol*, Vol. 27, PP.405-409, 2002.
- [12] KJ McElwee, DJ Tobin, JC Bystryn , Alopecia areata: an autoimmune disease, *Exp Dermatol*, Vol.8,PP. 371-379,1999.
- [13] T.Nakasa, S.Miyaki, A.Okubo, Expression of microRNA-146 in rheumatoid arthritis synovial tissue, *Arthritis Rheum*,Vol. 58,PP. 1284–1292, 2008.
- [14] R.M.O'Connell, D.S.Rao and D .Baltimore, MicroRNA regulation of inflammatory responses, *Annu Rev Immunol*, Vol.30, PP. 295-312, 2012.
- [15] K.M.Pauley, C.M.Stewart, A.E. Gauna, Altered miR-146a expression in Sjogren's syndrome and its functional role in innate immunity, *Eur J Immunol*,Vol.41,PP.2029-2039, 2011.

- [16] K.M.Pauley, S.Cha and E.K.Chan, MicroRNA in autoimmunity and autoimmune diseases. *J Autoimmun*, Vol. 32,PP.189–194, 2009.
- [17] C.H.Pratt, J.r.King.LE, A.G. Messenger, Alopecia areata,*Nat Rev Dis Primers*,Vol. 3,pp,17011, 2017.
- [18] F.Rajabi, L.A.Drake, M.M.Senna, Alopecia areata: a review of disease pathogenesis, *British J Dermatol*,Vol.179,PP.1033–1048, 2018.
- [19] T.A.Rodriguez, K.E.Fernandes, K.L.Dresser and M.Duvic, Concordance rate of alopecia areata in identical twins supports both genetic and environmental factors, *J Am Acad Dermatol* Vol. 62,PP.525–527, 2010
- [20] K.Smigielska-Czepiel , A.van den Berg, P.Jellema, Comprehensive analysis of miRNA expression in T-cell subsets of rheumatoid arthritis patients reveals defined signatures of naive and memory Tregs, *Genes Immun*,Vol.15 PP.115-125, 2014.
- [21] A Tahamtan, M Teymoori-Rad, B Nakstad, Anti-Inflammatory MicroRNAs and Their Potential for Inflammatory Diseases Treatment, *Front Immunol*,Vol. 9,PP.13-77, 2018.
- [22] Xu.Wang-Dong, Lu.Man-Man, Pan .Hai-Feng, Association of MicroRNA-146a with Autoimmune Diseases. *Inflammation*, Vol.35 ,PP.1525-1529, 2012.
- [23] E.H.C.Wang, G.M.Destefano, A.V. Patel, Identification of differentially expressed miRNAs in alopecia areata that target immune-regulatory pathways. *Genes Immunity*,Vol.8,PP.100-107, 2017.
- [24] G.Wang, L.S.Tam, E.K.Li., Serum and urinary cell-free MiR-146a and MiR-155 in patients with systemic lupus erythematosus, *J Rheumatol*,Vol.37,PP.2516–2522.2010.
- [25] H.Wei , M. Guan , Y. Qin , C.Xie , X.Fu , F.Gao , Circulating levels of miRNA 146a and IL-17 are significantly correlated with the clinical activity of Graves' ophthalmopathy, *Endocr J* , Vol.61,PP.1087-1092, 2014.
- [26] L.Xing, Z.Dai, A.Jabbari, Alopecia areata is driven by cytotoxic T lymphocytes and is reversed by JAK inhibition. *Nat Med*,Vol.20,PP.1043-1049, 2014.