

The Levels of Wnt5a and Its Receptors Frizzled5 and Frizzled2 as Immunohistochemical Biomarkers of Severity of Psoriasis

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Purpose: Psoriasis is a systemic, chronic and inflammatory condition. The exact pathogenesis is unclear. The abnormal expression of Wnt5a pathway in psoriasis vulgaris has been confirmed. Whether it is related to the severity of psoriasis is unclear.

Methods: Thirty-eight skin lesions from psoriasis vulgaris patients and 22 healthy adult skin tissues were taken. The semi-quantitative immunohistochemistry score of Wnt5a, Frizzled5 and Frizzled2 was evaluated under a microscope by two independent dermatologists. Psoriasis area and severity index (PASI) score system was used to evaluate the disease severity.

Results: The average PASI score of the patients was 16.25 ± 7.8 , and the average duration of disease was 19.6 ± 10.4 months. Wnt5a, Frizzled5 and Frizzled2 were highly expressed in psoriasis lesions. The semi-quantitative immunohistochemistry scores of Wnt5a, Frizzled5 and Frizzled2 were positively correlated with PASI scores ($r = 0.71$, $r = 0.46$, $r = 0.65$, respectively, all P -value < 0.01), but not correlated with duration of disease ($r = 0.11$, $r = 0.17$, $r = 0.29$, respectively, all P -value > 0.05). There were significant positive correlations between Wnt5a and Frizzled5 ($r = 0.57$, P -value < 0.01), as well as Wnt5a and Frizzled2 ($r = 0.59$, P -value < 0.01).

Conclusion: Wnt5a and its receptors play an important role in pathogenesis of psoriasis vulgaris and are positively correlated with the severity of psoriasis, and may be one of the immunohistochemical predictors of the severity of the disease.

Keywords: psoriasis vulgaris, Wnt5a, PASI, Frizzled5, Frizzled2

Introduction

Psoriasis is a chronic and systemic inflammatory condition with a prevalence is approximately 2–3%, and it may affect more than 125 million individuals worldwide.^{1,2} Its related mortality, economic burden and morbidity are high, and impairing the quality-of-life of psoriasis patients markedly. Excessive proliferation of keratinocytes, and infiltration of immune cells into the skin is involved in psoriasis, but the exact pathogenesis is unclear.³ The crucial role of Wnt5a in psoriasis had been investigated.^{4–6}

The Wnts are a large family with cysteine-rich secreted glycoproteins, which play an important role in the development and physiological processes of many tissues.⁷ As a key factor in the non-classical Wnt pathways, Wnt5a regulates the inflammatory response and various functions of the cells by activating a cascade signaling pathways, including cell proliferation, differentiation, apoptosis, survival, and cell polarity.⁸ Studies have revealed that Wnt5a gene was up-regulated in psoriatic lesions.^{4–6}

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Wnt5a signaling pathway can regulate downstream factors by binding related receptors, such as Frizzled5 and Frizzled2, and then to regulate cell biological function by activating β -catenin and protein kinase C (PKC). Wnt5a signaling pathway can regulate the proliferation, apoptosis and migration of keratinocytes.⁹ A recent analysis of gene pathogenicity found that Wnt5a gene is one of the five key genes and may be a biomarker for the diagnosis and treatment of psoriasis.¹⁰ It was also found in our previous studies that Wnt5a and its receptors were highly expressed in psoriatic lesions.¹¹

So we speculate that Wnt5a pathway is not only associated with psoriasis recurrence but also with psoriasis severity. The present study was planned to evaluate the correlation between Wnt5a, Frizzled5, Frizzled2 and the severity of the psoriasis.

Materials and Methods

Samples

Patients with psoriasis vulgaris with typical clinical manifestations and confirmed by histopathology were selected as research objects. Healthy adults with normal skin as the control group. All selected patients and healthy adults had no history of drug intake (such as glucocorticoids, retinoic acid, and immunosuppressive drugs) in the past 2 months, and used no any topical drugs in the past 2 weeks. There were no history of hypertension, diabetes, obesity and other chronic diseases when patients and healthy adults were included. All participants completed the written informed consent. All biopsies were done under local anesthesia. Evaluation of Psoriasis Area and Severity Index (PASI) scores was done by dermatologist. All experimental protocols were approved by the Institutional Review Board of the Xi'an Jiaotong University, and performed according to guidelines governing ethics care in China. This study was performed in accordance with the rules laid down in the Declaration of Helsinki.

Immunohistochemistry

All tissues were embedded in paraffin. At the temperature at 4°C, the paraffin sections were incubated with the primary anti-Wnt5a (Abcam ab110073, Cambridge, UK, final dilution 1:250), anti-Frizzled-2 (Abcam ab109094, Cambridge, UK, final dilution 1:300), and anti-Frizzled-5 (Abcam ab75234, Cambridge, UK, final dilution 1:250) antibody overnight. Then, a biocatalyst streptavidin-horseradish peroxidase-conjugated as secondary antibody (ZSGB-BIO, Beijing, China) were used, at last with hematoxylin counterstaining and 3'-diaminobenzidine (ZSGB-BIO, Beijing,

China). In negative control, phosphate buffered saline (PBS) was used to replace the primary antibody.

Semi-Quantitative Analysis of Immunohistochemistry Results

NanoZoomer Digital Pathology (Hamamatsu, Japan) was used to examine the sections. At high-power fields (200 \times magnifications), randomly selected five images were captured by a DP 70 CCD camera (Olympus Corp., Tokyo, Japan) for quantitative analysis. Semi-quantitative analysis of immunohistochemical results: the sample was scored with a percentage of positive cells (stain was cytoplasmic or nuclear) and a staining intensity score.¹² The staining intensity score was 0 for unstained, 1 for pale yellow, 2 for brown yellow, and 3 for brown. According to the percentage of the number of positive cells, the score was divided as: 0 points for <5%, 1 point for 5% to 25%, 2 points for 26% to 50%, and 3 points for >50%. Multiply the degree of staining and the percentage of stained cells in each section to get the final score: 0 to 1 is considered negative, and ≥ 2 is considered positive.

Statistical Analysis

All data were processed with SPSS19.0 statistical analysis software (SPSS Inc., Chicago, IL, USA). Unpaired *T* test, Chi-square test was used for statistical analysis of the results. Correlation analysis was performed by the Spearman two-variable correlation analysis method. $P < 0.05$ was defined as statistically differences.

Results

The Characteristics of the Participants

As shown in Table 1, 38 lesions from 38 patients with psoriasis vulgaris were enrolled (20 males and 18 females, age range 19–71 years, mean age was 50.2 ± 13.1 years). There were 16 lesions located at the trunk, 12 at the upper limb and 10 at the lower limb. The average PASI score was 16.25 ± 7.8 , and the average duration of disease was 19.6 ± 10.4 months. Normal skin tissue of 22 healthy adults (11 males and 11 females, age range 21–68 years,

Table 1 Clinical Data of the Study Participants

Clinical Data	Patients
Number of patients	38
Age (years), mean \pm SD	50.2 \pm 13.1
PASI score, mean \pm SD	16.25 \pm 7.8
Duration of psoriasis (months), mean \pm SD	19.6 \pm 10.4

Abbreviations: PASI, psoriasis area and severity index; SD, standard deviation.

Table 2 The Expression of Wnt5a, Frizzled5 and Frizzled2 in Psoriasis

Groups	N	Wnt5a		Frizzled5		Frizzled2	
		-	+	-	+	-	+
Psoriasis lesions	38	8(21.1%)	30(78.9%)	10(26.3%)	28(73.7%)	7(18.4%)	31(81.6)
Healthy controls	22	18(81.8%)	4(18.2%)	17(77.3%)	5(22.7%)	15(68.2%)	7(31.8%)

mean age was 48.7 ± 14.9 years) as the control. There were no significant differences in gender and age between the two groups (P -value > 0.05).

Immunohistochemical Analysis of Wnt5a, Frizzled5, and Frizzled2 in Psoriasis

As shown in Table 2 and Figure 1A and B. Thirty specimens of psoriasis had increased Wnt5a staining, not only in all the epidermal layers but also in

inflammatory and endothelial cells in the dermis. In normal skin, 18 cases had no increased in Wnt5a staining, Wnt5a proteins normally staining in the basal layer, and 4 cases had increased Wnt5a staining (in addition to the basal layer, there was a small amount positive cells in the spinous layer). The expression rate of Wnt5a in psoriasis was significantly higher than it in normal skin, and the difference was statistically significant ($\chi^2=20.95$, P -value < 0.001).

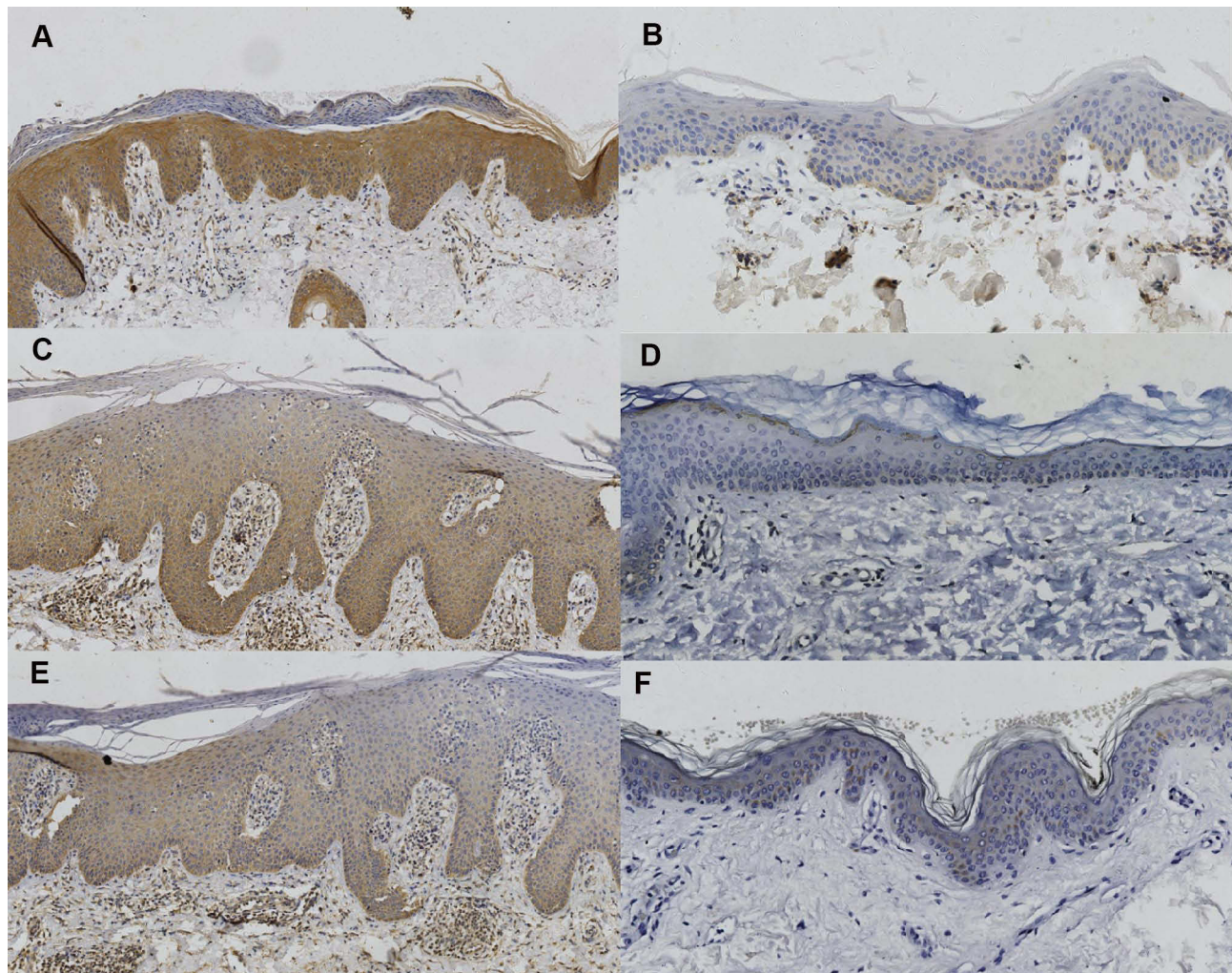


Figure 1 Skin sections immunostaining stained. (A) Wnt5a in lesions; (B) Wnt5a in normal skin; (C) Frizzled5 in lesions; (D) Frizzled5 in normal skin; (E) Frizzled2 in lesions; (F) Frizzled2 in normal skin. (A, C, E with $\times 100$ magnification, B, D, F with $\times 200$ magnification).

As shown in Table 2 and Figure 1C and D. There were 28 cases with psoriasis with increased of Frizzled5 staining in all the epidermal layers, inflammatory and endothelial cells in the dermis. In normal skin, 17 cases had normal of Frizzled5 staining (in the granular layer), and 5 cases had increased staining (there was a small amount positive cells in the spinous layer and the basal layer). The expression of Frizzled5 in psoriatic lesions was significantly higher than it in normal skin, and the difference was statistically significant ($\chi^2=12.63$, P -value <0.001).

As shown in Table 2 and Figure 1E and F, 31 psoriasis cases (81.58%) had increased Frizzled2 staining in all the epidermal layers, inflammatory and endothelial cells in dermis. In normal skin tissues, 7 cases had slight increased staining (besides the basal layer, there was increased staining in the upper basal layer). The expression of Frizzled2 in psoriatic tissue was significantly higher than it in normal skin, and the difference was statistically significant ($\chi^2 = 12.79$, P -value < 0.001).

Correlation Between the Expression Levels of Wnt5a and Frizzled5, Frizzled2 in Psoriasis

Spearman two-variable correlation coefficient analysis showed that significant positive correlations between Wnt5a and Frizzled5 ($r = 0.57$, P -value < 0.01), as well as Wnt5a and Frizzled2 ($r = 0.59$, P -value < 0.01) in psoriasis.

Correlation Between Semi-Quantitative Scores of Wnt5a, Frizzled5 and Frizzled2 with PASI Score and Duration of Disease

Semi-quantitative score of Wnt5a in psoriasis vulgaris lesions was positively correlated with PASI score ($r = 0.71$, P -value < 0.01), but not correlation with the duration of disease ($r = 0.11$, P -value > 0.05) (Figure 2A and B). There was a positive correlation between the semi-quantitative score of Frizzled5 and the PASI score ($r = 0.46$, P -value < 0.01), and there was no correlation between the semi-quantitative score of Frizzled5 and the duration of disease ($r = 0.17$, P -value > 0.05) (Figure 2C and D). Semi-quantitative score of Frizzled2 was positively correlated with PASI score ($r = 0.65$, P -value < 0.01), but not correlated with the duration of disease ($r = 0.29$, P -value > 0.05) (Figure 2E and F).

Discussion

In psoriatic lesions, Wnt5a and the receptors Frizzled2 and Frizzled5 were not only expressed in the entire epidermis but also strongly positive expressed in superficial dermal inflammatory cells and vascular endothelial cells, indicating that the Wnt5a pathways may be involved in the regulation of inflammatory response process, but its specific role in psoriasis is currently unknown.

Infiltration of inflammatory cells in the epidermis and dermis in psoriatic lesions is a crucial factor in the pathogenesis of psoriasis. Numerous factors have been found to

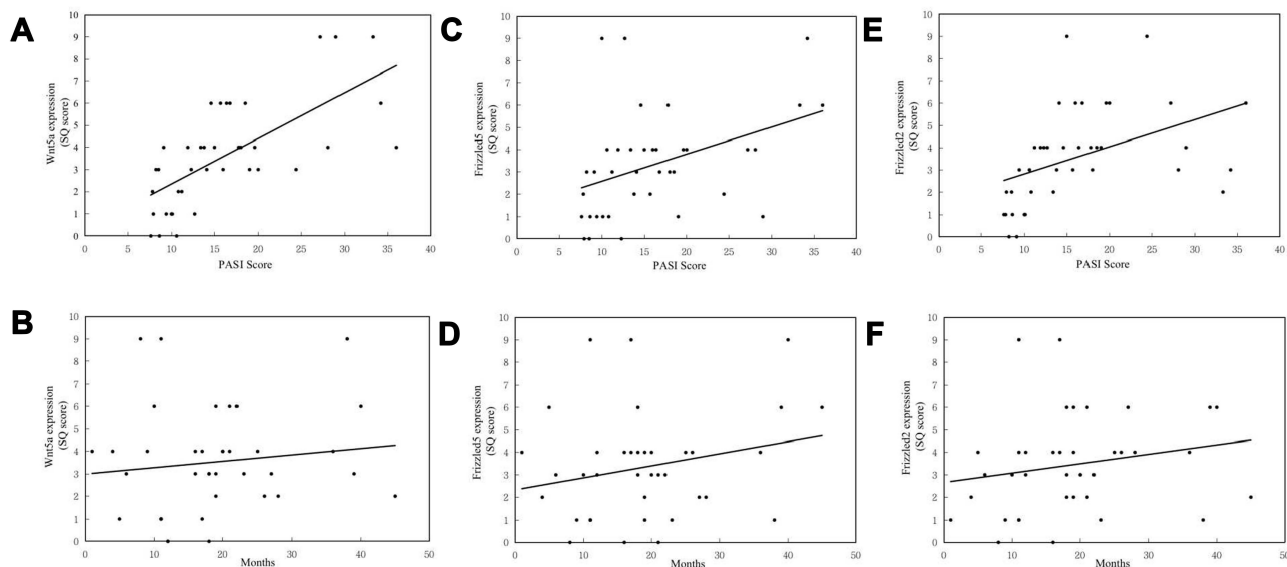


Figure 2 (A) The semi-quantitative score of Wnt5a in lesions was positively correlated with PASI score; (B) but not with duration of disease; (C) the semi-quantitative score of Frizzled5 was positively correlated with PASI score; (D) but not with duration of disease; (E) the semi-quantitative score of Frizzled2 was positively correlated with PASI score; (F) but not with duration of disease.

participate in the pathogenesis of psoriasis vulgaris in mouse models.¹³ The Wnt5a pathway plays an important role in the pathogenesis of various inflammatory diseases.¹⁴ After binding to the corresponding receptors, Wnt5a activate the downstream molecular pathways, and then affects the progress of the diseases. Interventions targeting Wnt5a or its receptors and downstream factors may be one of the methods for treating inflammatory diseases.¹⁵ Wnt5a and receptors are expressed in epithelial cells, lymphocytes, endothelial cells and smooth muscle cells.¹⁶ In HaCaT cells, after treatment with Wnt5a, the expression levels of inflammatory markers, including IFN- γ , IL-8 and IL-17A were unregulated.¹⁷ IL-1 α , TNF- α , IFN- γ , and TGF- α can induce the expression of Wnt5a.⁵ All these inflammatory markers play critical roles in the pathogenesis and maintenance of psoriasis.³ Wnt5a also participates in the early emergency response of psoriasis to mechanical stress.¹⁸ Differential methylations of Wnt5a and Frizzled2 were found in psoriasis.¹⁹ Nuclear factors of activated T-cells, cytoplasmic1 (NFATc1), a downstream gene of Wnt5a, can induce psoriasisform dermatitis.²⁰ Wnt5a-activated signalling pathways mediate interactions among keratinocytes, immune cells and inflammatory factors, and play an important role in the development of psoriasis and psoriatic arthritis (PsA).²¹ Therefore, the Wnt5a pathway may be a key factor in the inflammatory cytokines loop and enables the occurrence of cytokine storm of psoriasis.

Frizzled2 activates the classical β -catenin and the independent β -catenin pathway by regulating the calcium ion pathway.²² Wnt5a/Frizzled2 can activate small G protein, activated small G protein can activate Phospholipase C (PLC), PLC induces endoplasmic reticulum to release calcium ions, which further activate calcineurin or PKC pathway.²² This pathway is also used to treat psoriasis, eg, calcipotriol (a vitamin D3 derivative drug) inhibits psoriatic keratinocyte proliferation through PKC and PLC. Tacrolimus and cyclosporin may also inhibit the immune response of psoriasis by inhibiting phosphatase. Wnt5a/Frizzled5 pathway can induce the synthesis of TNF- α and IL-1, both TNF- α and IL-1 are involved in the pathogenesis of psoriasis.²³ Wnt5a/Frizzled5 is highly expressed in rheumatoid arthritis. Fibroblasts isolated from rheumatoid joints can synthesize IL-15, IL-8 and IL-6, and then these cytokines act on the Wnt5a/Frizzled5 signaling pathway.²⁴ In endothelial cells, Wnt5a induces Nuclear factor of activated T-cells (NFAT) transcription by binding to Frizzled5, and then induces T-cells to secrete IL-2, both NFAT and IL-2 are strongly

expressed in psoriasis vulgaris.²⁵ In our current research, Wnt5a was positively correlated with the expression of Frizzled2 and Frizzled5 in psoriasis lesions. Therefore, Wnt5a may be bind to Frizzled2 and Frizzled5, to activate related signaling pathways, and then regulate downstream factors to participate in the pathogenesis of psoriasis.

The correlation analysis found that the expression of Wnt5a, Frizzled2 and Frizzled5 were positively correlated with the severity of psoriasis vulgaris (PASI score), but not to the duration of the disease. The degree of erythema, scale and cell infiltration are important indicators for clinical evaluation of the severity of psoriasis. The Wnt and Wnt5a pathways may be as potential therapeutic targets of inflammatory disorders and skin cancer.^{26,27} Thus, with the study of the Wnt5a signaling pathway, it as a target for psoriasis treatment will become a reality in the future. This study also has some limitations, such as the expression of Wnt5a and its receptors Frizzled5 and Frizzled2 in skin lesions after treatment were not detected. The number of patients in this study is too small, and it needs to be further expanded, and the animal experiments need to be carried out in the future.

Conclusion

In summary, our results further strengthen the linkage between psoriasis and the Wnt5a signaling pathway. Wnt5a, Frizzled2 and Frizzled5 could be used as immunohistochemical biomarkers of severity of psoriasis.

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Disclosure

The authors declare no conflicts of interest in this work.

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