Investigating the **TLR9 mRNA** Expression Level in Different Histological Types of Colorectal Polyps

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**Abstract**

*TLR9* is a cellular DNA receptor of the innate immune system which plays a pivotal role in inflammatory response. Recently, changing expression levels of *TLR9* has been observed in a wide range of cancer cells; however, there is little information about colorectal polyps. Herein, we assessed the mRNA expression of *TLR9* in different colorectal polyp types compared to normal group in order to investigate its expression level during CRC initiation. Fifty-four biopsy samples from colorectal polyp patients and from 20 healthy subjects were collected. The mucosal mRNA expression level of *TLR9* gene was identified by real time PCR. Fold change of gene expression was evaluated by 2⁻ΔΔct method. There was a significant relationship between the lower expression of *TLR9* gene in the polyp cases compared to normal individuals (P value = 0.0005), Also, decreased *TLR9 mRNA* expression was obtained in adenomas in contrast to hyperplastic and normal groups (P value = 0.0008). Based on the current results, we hypothesized that aberrant surface expression of *TLR9* on tumor cells may promote the growth and invasion of colorectal polyps. Further, *TLR9* modulation may have an important impact on the development of novel therapeutic strategies.

**Keywords**: Colorectal Polyp- mRNA expression- *TLR9*

**Introduction**

Colorectal cancer (CRC) is a considerable global health problem in the world and the third diagnosed malignancy following lung and breast cancer (Peters et al., 2016; Hale et al., 2017). CRC is one of the major causes of cancer death in western countries and also the third most common cancer in Iran (Ansari et al., 2006; Haghdoost et al., 2011). Most CRC cases progress from precursors recognized as colorectal polyps; the two common types of polyps include: hyperplastic (HP) and adenomatous polyps (AP). AP itself is divided in three main sub-groups of tubular adenoma (TP), tubuvillous (TVP), and villus (VP). Previous investigations (Leslie et al., 2002; Kim et al., 2011) have shown that colorectal tumors may originate from transformed polyps within several years. Histologically, colorectal polyps are found in different shapes including adenoma, hyperplastic, hamartomatous, and inflammatory polyps (Leslie et al., 2002). Neoplastic or adenoma polyps are more important as they are suspected of being malignant and cancerous (O’Connell and Crockett, 2017). Indeed, recent investigations have revealed that hyperplastic polyps have the malignancy potential in some cases (Liljegren et al., 2003; Inoue et al., 2007). The relationship between inflammation and tumor is becoming increasingly important in the study on the pathogenesis of CRC. Several studies have confirmed that TLRs as immune molecules could mediate inflammatory response and play an important role in this process (Chen et al., 2007; Kolumam et al., 2005; Siegel et al., 2016). Nevertheless, the role of TLR9-mediated immune inflammation reaction in the process is not currently clear (Caixia et al., 2018). In this study, we dealt with evaluating the expression level of the *TLR9* in colonic AP and HP cases in contrast to healthy subjects and with finding the relationship between *TLR9 mRNA* expression level and pathological features of AP and HP cases.

**Materials and Methods**

**Human sample collection**

The present study was case control and the investigated population (54 patients with polyp and 20 normal individuals) was chosen from the cases with colorectal polyps who were referred to Taleghani hospital, Shahid Beheshti University of Medical Sciences, Tehran-Iran.

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between October 2015 and April 2017. The sample was chosen through ransom sampling. Colorectal polyps were identified through colonoscopy and confirmed by pathology results. Polyp-free controls were defined as those with no polyps identified during colonoscopy and no previous history of colorectal polyps. Polyp type classifications were done according to polyp histology: hyperplasic (HP), tubular adenoma (TA), and tubulovillous polyp (TVP), where dysplasia grades and polyp sizes were characterized by a pathologist. The clinical information of patients was collected by a questionnaire. The study was approved by the Clinical Research Ethics Committee of Shahid Beheshti University of Medical Sciences and the Ethics Committee of Taleghani Hospital, Tehran, Iran with No.2014/770.

RNA extraction and quality control

Total RNA was extracted from all samples (Yekta Tajhiz Azma kit, Teheran, Iran) according to the kit instructions. RNA concentration and purity ratios (OD260/280, OD260/230) were evaluated by NanoDrop 1000 spectrophotometer (NanoDrop Technologies, Wilmington, DE, USA). The integrity of RNA was determined by electrophoresis on a denaturing 1.5% agarose gel.

cDNA synthesis

Total RNAs were converted to cDNA by Retrotransciptase (RT) reaction (TaKaRa kit, Cat No.RR037A, Otsu, Shiga, Japan) according to the following: 2 mg of total RNA was picked up and denatured at 95°C for 5 min. Thereafter, the tubes were placed on ice where 5 μL of 5× primer script buffer, 0.5 μL RT enzyme, 1.24 μM oligo dt primer, 10 μM random 6 mer, 1 μM ribolock, 1 μL easy dilution, and 5 μL RNA free distilled water (dH2O), were added. The cDNA synthesis was performed as follows: 25°C for 5 min, 42°C for 15 min, 85°C for 1 min for inactivation of the reverse transcriptase enzyme and 4°C for 10 min for hold temperature. Next, cDNA products were stored at -20°C. Note that in all reactions, the same concentrations of RNA samples were used (RNA adjustment).

Real time-PCR

Real-time PCR was performed following the standard SYBR Premix Ex Taq™ kit (TaKaRa Bio Inc., Otsu, Japan) protocol, using a final volume of 20 μL containing 5 μL of reverse-transcribed cDNA and 1 μL of 10 pmol forward and reverse primers, using Applied Biosystems 7500 version 1 (ABI, Foster City, CA, USA). Real-time PCR was carried out by expressive primers (Table 1) under the following conditions: 95°C for 5 s, 40 cycles of 95°C for 5 s, 60°C for 34 s, 95°C for 15 s, 60°C for 1 s and 60°C for 15s (Khatibi et al., 2018). Amplification signals for samples were normalized by glyceraldehyde 3-phosphate dehydrogenase (GAPDH) gene. Fold change of gene expression was evaluated by $2^{-ΔΔct}$ method.

Statistical analysis

All results were analyzed by Graph pad Prism software version 5 (San Diego, CA, USA). The data were non-normally distributed and the non-parametric test was used. Specifically, student t-test and one-way ANOVA test were performed. $P$-value < 0.05 was considered statistically significant.

Results

Among 54 polyp cases, 26 (48.3%) were identified as hyperplastic, 7 (12.9%) as tubular, and 21 (38.8%) as tubulovillous. Clinicopathological parameters of patients with different polyp types are presented in Table 2. Based on these results, there was a significant relationship between the lower expression level of TLR9 in the polyp cases compared with normal samples ($P$-value = 0.0005) (Figure 1). Also, comparing AP (TP, TVP), HP, and

![Figure 1. The Significant Down-Regulation of TLR9 mRNA in the Polyp Group Compared with the Normal Group ($P$-value = 0.0005)](image)

Table 2. Clinicopathological Parameters of Patient with Different Polyps

| Parameter | Type     | Number |
|-----------|----------|--------|
| Polyp types | Hyperplastic | 26 (48.3%) |
|            | Tubular   | 7 (12.9%) |
|            | Tubulovillous | 21 (38.8%) |
| Dysplasia  | HGD      | 24 (44.5%) |
|            | LGD      | 30 (55.5%) |
| Gender     | Male     | 28 (52%) |
|            | Female   | 26 (48%) |
| Site       | Rectum   | 16 (29.6%) |
|            | AC       | 7 (13%) |
|            | AD       | 15 (27.8%) |
|            | TC       | 16 (29.6%) |
| Total      |          | 54 (100%) |

Table 1. Oligonucleotides Pair of Primers Used for the Relative Gene Expression

| Gene | Primer Sequence |
|------|-----------------|
| TLR9-F | 5’_AATCCCTCATATCCCTGTCCC_3’ |
| TLR9-R | 5’_GTTGCCGTCCATGAATAGGAAG_3’ |
| GAPDH-F | 5’_TGACTTTCAACAGCGACACCCA-3’ |
| GAPDH-R | 5’_CACCTGTTGCTGTAGCCAAA-3’ |
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LGD compared with normal participants. Also, we found various levels of TLR9 mRNA in different polyp types such as TP and TVP types. Our results were in contrast with Yesudhas et al., (2014) indicating that TLR7, 8, 9, and 10 mRNA expression increased in tissues of CRC patients. Indeed, they worked on CRC cases while the present study focused on different polyp cases; hence these differences may arise from differences between polyp and tumor tissues. On the other hand, Sandholm et al., (2014) demonstrated the beneficial role of TLR9 on breast cancer treatment (Sandholm and Selander, 2014). Also, Shahriari et al., (2017) hypothesized that aberrant surface expression of TLR9 on tumor cells may promote tumor growth and invasion. They highlighted a dual contradictory role for CpG-ODNs, as an adjuvant agent in cancer therapy. Hasan et al., (2007) as well as Pacini et al., (2015) reported that downregulation of TLR9 may be a crucial step in the carcinogenic events of human papillomavirus associated with cervical cancer. Furthermore, some researchers declared that oncogenic viruses such as papillomavirus 16, Epstein-Barr virus, and hepatitis B virus may have an important role in downregulating TLR9 expression (Fathallah et al., 2010; van Gent et al., 2011; Vincent et al. 2011). Finally, Jouhi et al., (2015) findings revealed diminishing expression levels of TLR9 during Merkel cell carcinoma.

In conclusion, we observed decreasing TLR9 expression in polyp cases of patients compared to normal samples. Also, lower expression level of TLR9 in HGD and LGD compared to normal participants was achieved and different expression levels were obtained between TP and TVP. According to these results, it can be concluded that the reduction of TLR9 expression may play a significant role in the progression of polyps to CRC and malignancy. Indeed, cancer cells may be able to escape the immune system due to TLR9 reduction. Finally, TLR9 modulation may have an impact on the development of novel therapeutic strategies.

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