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# ACCEPTED MANUSCRIPT

#### Distinctive microRNA signature associated of neoplasms with the

## Wnt/β-catenin signaling pathway

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**Abbreviation**: FZD, Frizzled; LDL receptor–related protein, LRP; Dvl, Dishevelled; APC, adenomatous polyposis coli; CK1 $\alpha$ , casein kinase 1 $\alpha$ ; GSK3 $\beta$ , glycogen synthase kinase 3 $\beta$ ; HCC, hepatocellular carcinoma; EZH2, zeste homologue 2; sFRPs, secreted Frizzled-related proteins; DKK, Dickkop; miRs, MicroRNAs; WIF, Wnt inhibitory factor; TCF, T-cell factor; LEF, lymphoid enhancer-binding factor; IRS, insulin receptor substrate; NLK, Nemo-like kinase.

## ABSTRACT

As the crucial biological regulators, microRNAs that act by suppressing their target genes are involved in a variety of pathophysiological processes. It is generally accepted that microRNAs are often dysregulated in many types of neoplasm and other human diseases. In neoplasm, microRNAs may function as oncogenes or tumor suppressors. As constitutive activation of the Wnt signaling pathway is a common feature of neoplasm and contributes to its development, progression and metastasis in various cancers, numerous studies have revealed that microRNAs-mediated gene regulation are interconnected with the Wnt/ $\beta$ -catenin signaling pathway, forming an

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