

ISSN: 2574 -1241 DOI: 10.26717/BJSTR.2025.63.009830

Shifting Correlative Trends of G-C% Content and 3'UTR Length

Shubh R Nahar¹, Md Mostafizur Rahman¹, Franziska Ahrend^{1,2}, Travis Varnum³, Giulianna Rivero³, Morgan Seymour³, Mary Katherine Swanson Lee³, Syeda Saberin Jahan^{4,7}, Kai Ding⁵, Bill Zhong¹, Long Ma¹, Valerii Drozhenko⁷ and Alexander Kofman^{6,7*}

ARTICLE INFO

Received: August 12, 2025

Published: August 21, 2025

Citation: Shubh R Nahar, Md Mostafizur Rahman, Franziska Ahrend, Travis Varnum, Giulianna Rivero, Morgan Seymour, Mary Katherine Swanson Lee, Syeda Saberin Jahan, Kai Ding, Bill Zhong, Long Ma, Valerii Drozhenko and Alexander Kofman. Shifting Correlative Trends of G-C% Content and 3'UTR Length. Biomed J Sci & Tech Res 63(1)-2025. BJSTR. MS.ID.009830.

ABSTRACT

We report the moderate positive correlation (r=0.52), between low G-C% content and the length of the 3'UTRs, as well as moderate negative correlation (r=-0.57) between high G-C% content and the 3'UTR length. The results corroborate our previous data on the asymmetric distribution of microRNA target sites along the 3'UTR and point to the possible influence of nucleotide composition on the cis-acting regulatory elements within the 3'UTR.

Keywords: mRNA; 3'UTR; Nucleotide

Abbreviations: mRNA: messenger RNA; 3UTR: Untranslated Region

Short Communication

The growing number of reports indicate that codon usage bias and nucleotide content are among the important regulatory factors influencing gene expression at various levels. They may impact the nuclear mRNA concentration [1] and mRNA stability [2,3] and alter the expression of the heterologous genes in bacteria, plants, yeast, mammalian cells, and transgenic animals [4]. Previously we report-

ed the moderate negative correlation between the 3'UTR length and the G-C% content in the 3'UTRs of 38 human mRNAs (in press). In this study we used the larger dataset of the prevailing transcripts of 824 human genes (RefSeq database [5]). We found the moderate positive correlation (r=0.52), between the low G-C% content (less than 30%) and the length of the 3'UTRs, as well as moderate negative correlation (r=-0.57) between the high G-C% content (more than 60%)

¹Department of Computer Science, Troy University, USA

²National Institute of Diabetes and Digestive and Kidney Diseases, National Institutes of Health, USA

³Department of Biological and Environmental Sciences, Troy University, USA

⁴Jalalabad Ragib-Rabeya Medical College and Hospital, Bangladesh

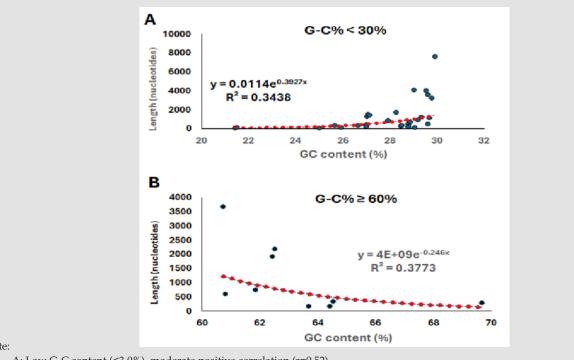
⁵John Hopkins University Medical School, USA

⁶University of West Alabama, USA

⁷Filatov Institute of Eye Diseases and Tissue Therapy, the National Academy of Medical Sciences, Odesa, Ukraine

^{*}Corresponding author: Alexander Kofman, Human Biology Variations Computational Genomics Research, Marazliivska St. 2, Odessa, Ukraine

and the 3'UTR length (Figure 1). The results indicate some similarity with the previously obtained data on asymmetric distribution of microRNA-target sites along the 3'UTR and suggest a complex interplay between the nucleotide composition and the 3'UTR regulatory functions, which remain to be elucidated.



Note:

- A: Low G-C content (<3 0%), moderate positive correlation (r=0.52).
- B: High G-C content (\geq 60%), moderate negative correlation (r=-0.57).

Figure 1: Moderate correlation between the 3'UTR length (human mRNAs) and G-C% content.

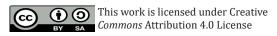
References

- Qian W, Zhang J (2021) Codon usage bias and nuclear mRNA concentration: Correlation vs. causation. Proc Natl Acad Sci 118.
- Mishima Y, Tomari Y (2016) Codon Usage and 3' UTR Length Determine Maternal mRNA Stability in Zebrafish. Mol Cell 61: 874-885.
- Hanson G, Coller J (2018) Codon optimality, bias and usage in translation and mRNA decay. Nat Rev Mol Cell Biol 19: 20-30.
- 4. Kofman Alexander, Marcus Graf, Alexandra Bojak, Ludwig Deml, Kurt Bieler, et al. (2003) HIV-1 gag expression is quantitatively dependent on the ratio of native and optimized codons. Tsitologiia 45: 86-93.
- O Leary Nuala A, Mathew W Wright, J Rodney Brister, Stacy Ciufo, Diana Haddad, et al. (2016) Reference sequence (RefSeq) database at NCBI: current status, taxonomic expansion, and functional annotation. Nucleic Acids Res 44: 733-745.

ISSN: 2574-1241

DOI: 10.26717/BISTR.2025.63.009830

Alexander Kofman. Biomed J Sci & Tech Res



Submission Link: https://biomedres.us/submit-manuscript.php



Assets of Publishing with us

- Global archiving of articles
- Immediate, unrestricted online access
- Rigorous Peer Review Process
- Authors Retain Copyrights
- Unique DOI for all articles

https://biomedres.us/