



OPEN ACCESS

EDITED AND REVIEWED BY

Nicoletta Potenza,
University of Campania Luigi Vanvitelli, Italy

*CORRESPONDENCE

Lingling Wang,
✉ wl_198927@126.com
Bin Hu,
✉ binhu0612@foxmail.com
Min-Jin Han,
✉ minjinhan@126.com
Q.-Z. Zhou,
✉ cnzqzhou@foxmail.com

RECEIVED 10 January 2025

ACCEPTED 13 January 2025

PUBLISHED 07 February 2025

CITATION

Wang L, Hu B, Han M-J and Zhou Q-Z (2025)
Editorial: The non-coding RNA world in animals
and plants.
Front. Genet. 16:1558406.
doi: 10.3389/fgene.2025.1558406

COPYRIGHT

© 2025 Wang, Hu, Han and Zhou. This is an open-access article distributed under the terms of the [Creative Commons Attribution License \(CC BY\)](#). The use, distribution or reproduction in other forums is permitted, provided the original author(s) and the copyright owner(s) are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.

Editorial: The non-coding RNA world in animals and plants

Lingling Wang^{1*}, Bin Hu^{2*}, Min-Jin Han^{3,4*} and Q.-Z. Zhou^{5*}

¹Ministry of Education Key Laboratory for Ecology of Tropical Islands, Key Laboratory of Tropical Animal and Plant Ecology of Hainan Province, College of Life Sciences, Hainan Normal University, Haikou, China, ²Rubber Research Institute, Chinese Academy of Tropical Agricultural Sciences, Haikou, China, ³State Key Laboratory of Resource Insects, Institute of Sericulture and Systems Biology, Southwest University, Chongqing, China, ⁴Key Laboratory of Sericultural Biology and Genetic Breeding, Ministry of Agriculture and Rural Affairs, College of Sericulture, Textile and Biomass Sciences, Southwest University, Chongqing, China, ⁵DUKE-NUS Medical School, National University of Singapore, Singapore, Singapore

KEYWORDS

non-coding RNAs, long noncoding RNA, microRNA, plant, animal

Editorial on the Research Topic

The non-coding RNA world in animals and plants

Non-coding RNAs (ncRNAs) are transcriptionally produced and do not code proteins, with a spectrum of diversified type and function (Chen and Kim, 2024). Although ncRNAs have many types, including the long noncoding RNAs (lncRNAs), microRNAs (miRNAs), small nucleolar RNAs (snoRNAs), transfer RNA (tRNA), ribosomal RNA (rRNA), and circular RNA (circRNA), most of them are lncRNAs and miRNAs. In 1984, the first regulatory ncRNA *microRNA F (micF)* was discovered from *Escherichia coli*. *MicF* inhibits the translation of *OmpF* by binding to its ribosome-binding site, blocking ribosomal entry (Mizuno et al., 1984; Corcoran et al., 2012).

Over the past 40 years, numerous regulatory ncRNAs, such as *Let-7*, *enod40*, *H19*, and *NEAT1*, have been identified as key regulators in animals or plants (Bartolomei et al., 1991; Crespi et al., 1994; Pasquinelli et al., 2000; Hutchinson et al., 2007). Studies have demonstrated the crucial roles of ncRNAs—particularly miRNAs and lncRNAs—in development, metabolism, and disease (Kallen et al., 2013; Tripathi et al., 2013; Chen and Kim, 2024). However, most of the ncRNA world is still unknown.

To provide new insights and a deeper understanding of the functional and regulatory roles of ncRNAs, we curated six articles for the “*The Non-Coding RNA World in Animals and Plants*” topic. These papers include original research on ncRNAs in tobacco, common carp (*Cyprinus carpio* L.), and pulmonary hypertension, as well as reviews on ncRNAs in endocrine disorders, eye diseases, and intrahepatic cholestasis of pregnancy (ICP).

Xie et al. analyzed 549 publicly available RNA-seq datasets on tobacco and identified 30,212 lncRNAs (Xie et al.). These lncRNAs exhibit distinct characteristics compared to coding genes, including fewer exons, higher A/U content, and greater tissue specificity. Functional analysis of the potential targets of these lncRNAs revealed their association with nicotine biosynthesis; their findings further validated through topping treatment. This study advances our understanding of the functional roles of lncRNAs in tobacco and provides new candidate genes for regulating nicotine production.

Ledesma-Pacheco et al. presented our current understanding regarding miRNAs' regulatory mechanism of endocrine disorders and their potential influence as disease biomarkers during their development processes by overviewing recent and significant research outputs (Ledesma-Pacheco et al.). This review illuminated the most recent

information related to the potential functions of miRNAs in endocrine disorders, including diabetes mellitus, thyroid diseases, and osteoporosis, and their novel diagnostic and therapeutic purposes. It will enhance our knowledge of miRNAs' roles in endocrine disorders and facilitate the development of novel miRNA-based diagnostic and therapeutic tools for studying endocrine disorders.

Benavides-Aguilar et al. integrated recent studies on miRNAs in common eye diseases to illustrate the regulatory roles of miRNAs in the eye-related diseases such as cataracts, glaucoma, and macular degeneration (Benavides-Aguilar et al.). This review provides valuable insights into the potential applications of miRNAs in the prognosis and treatment of these eye-related disorders.

Das et al. integrated 468 raw RNA-seq datasets from 28 tissues in common carp (*C. carpio* L.), a substitute vertebrate fish model for zebrafish, to identify lncRNAs and circular RNAs using various bioinformatics tools (Das et al.). They conducted the lncRNA-miRNA-mRNA interaction network analysis and introduced CCncRNadb, a comprehensive web resource (<http://backlin.cabgrid.res.in/ccncrnadb/>) to facilitate further exploration of ncRNAs in common carp.

Xiong et al. consolidated recent discoveries on miRNAs, lncRNAs, circRNAs, etc. for ICP to discuss their potential as diagnostic markers, prognostic tools, and therapeutic targets, offering a foundation for improving early detection and personalized treatment of ICP (Xiong et al.).

Chen et al. conducted a comprehensive study on the peripheral blood samples of patients with pulmonary hypertension (PH) and healthy individuals through whole-genome miRNA sequencing and transcriptome analysis, exploring the potential role of miRNAs in PH (Chen et al.). By screening the differentially expressed miRNAs, they identified four miRNAs (hsa-miR-1304-3p, hsa-miR-490-3p, hsa-miR-11400, and hsa-miR-31-5p) as potential clinical diagnostic biomarkers for PH. This finding provides a valuable foundation for further understanding the specific role of miRNAs in the mechanisms of PH, opening new avenues for early diagnosis and precision medicine in PH.

In summary, these articles expand our current knowledge of ncRNAs and deepen our understanding of the non-coding RNA world.

References

- Bartolomei, M. S., Zemel, S., and Tilghman, S. M. (1991). Parental imprinting of the mouse *H19* gene. *Nature* 351, 153–155. doi:10.1038/351153a0
- Chen, L. L., and Kim, V. N. (2024). Small and long non-coding RNAs: past, present, and future. *Cell* 187, 6451–6485. doi:10.1016/j.cell.2024.10.024
- Corcoran, C. P., Podkaminski, D., Papenfort, K., Urban, J. H., Hinton, J. C., and Vogel, J. (2012). Superfolder GFP reporters validate diverse new mRNA targets of the classic porin regulator, MicF RNA. *Mol. Microbiol.* 84, 428–445. doi:10.1111/j.1365-2958.2012.08031.x
- Crespi, M. D., Jurkevitch, E., Poirer, M., D'aubenton-Carafa, Y., Petrovics, G., Kondorosi, E., et al. (1994). *enod40*, a gene expressed during nodule organogenesis, codes for a non-translatable RNA involved in plant growth. *EMBO J.* 13, 5099–5112. doi:10.1002/j.1460-2075.1994.tb06839.x
- Hutchinson, J. N., Ensminger, A. W., Clemson, C. M., Lynch, C. R., Lawrence, J. B., and Chess, A. (2007). A screen for nuclear transcripts identifies two linked noncoding RNAs associated with SC35 splicing domains. *BMC Genomics* 8, 39. doi:10.1186/1471-2164-8-39
- Kallen, A. N., Zhou, X. B., Xu, J., Qiao, C., Ma, J., Yan, L., et al. (2013). The imprinted *H19* lncRNA antagonizes *let-7* microRNAs. *Mol. Cell* 52, 101–112. doi:10.1016/j.molcel.2013.08.027
- Mizuno, T., Chou, M. Y., and Inouye, M. (1984). A unique mechanism regulating gene expression: translational inhibition by a complementary RNA transcript (micRNA). *Proc. Natl. Acad. Sci. U. S. A.* 81, 1966–1970. doi:10.1073/pnas.81.7.1966
- Pasquinelli, A. E., Reinhart, B. J., Slack, F., Martindale, M. Q., Kuroda, M. I., Maller, B., et al. (2000). Conservation of the sequence and temporal expression of *let-7* heterochronic regulatory RNA. *Nature* 408, 86–89. doi:10.1038/35040556
- Tripathi, V., Shen, Z., Chakraborty, A., Giri, S., Freier, S. M., Wu, X., et al. (2013). Long noncoding RNA *MALAT1* controls cell cycle progression by regulating the expression of oncogenic transcription factor B-MYB. *PLoS Genet.* 9, e1003368. doi:10.1371/journal.pgen.1003368

Author contributions

LW: writing–original draft and writing–review and editing. BH: writing–original draft and writing–review and editing. M-JH: writing–original draft and writing–review and editing. Q-ZZ: writing–original draft and writing–review and editing.

Funding

The author(s) declare that no financial support was received for the research, authorship, and/or publication of this article.

Acknowledgments

The authors would like thank the reviewers and authors for their contribution to this Research Topic.

Conflict of interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Generative AI statement

The author(s) declare that no Generative AI was used in the creation of this manuscript.

Publisher's note

All claims expressed in this article are solely those of the authors and do not necessarily represent those of their affiliated organizations, or those of the publisher, the editors and the reviewers. Any product that may be evaluated in this article, or claim that may be made by its manufacturer, is not guaranteed or endorsed by the publisher.