李振豪醫師論文與著作

1. *	Chung-Lin Lee*, **Chen-Hao Lee***, Chih-Kuang Chuang, Huei-Ching Chiu, Yen-Jiun Chen, Chao-Ling Chou, Peih-Shan Wu, Chih-Ping Chen, Hsiang-Yu Lin, Shuan-Pei Lin. Array-CGH increased the diagnostic rate of developmental delay or intellectual disability in Taiwan. Pediatrics and Neonatology, 2019 Aug;60(4):453-460
2.	En-Ting Wu, Wuh-Liang Hwu, Yin-Hsiu Chien, Ching Hsu, Ting-Fu Chen, Nai-Qi Chen, Hung-Chieh Chou, Po-Nien Tsao, Pi-Chuan Fan, I-Jung Tsai, Shuan-Pei Lin, Wu-Shiun Hsieh, Tung-Ming Chang, Chi-Nien Chen, **Chen-Hao Lee**, Yen-Yin Chou, Pao-Chin Chiu, Wen-Hui Tsai, Hann-Chang Hsiung, Feipei Lai, Ni-Chung Lee. Critical Trio Exome Benefits In-Time Decision-Making for Pediatric Patients with Severe Illnesses. Pediatric Critical Care Medicine, 2019 Nov;20(11):1021-1026.
3.	Yin-Hsiu Chien, Ni-Chung Lee, Pin-Wen Chen, Hui-Ying Yeh, Michael H Gelb, Pao-Chin Chiu, Shao-Yin Chu, **Chen-Hao Lee**, An-Ru Lee, Wuh-Liang Hwu. Newborn screening for Morquio disease and other lysosomal storage diseases: results from the 8-plex assay for 70,000 newborns. Orphanet Journal of Rare Diseases, 2020 Feb 3;15(1):38.
4.	Wen-Li Lu, Po-Chih Shen, **Chen-Hao Lee**, Yu-Tsun Su, Li-Min Chen. High Risk of Early Cataracts in Young Type 1 Diabetes Group: A Nationwide Cohort Study. International Journal of Endocrinology, 2020 Oct 9;2020:8160256.
5.	Hsiang-Yu Lin, Chung-Lin Lee, Sisca Fran, Ru-Yi Tu, Ya-Hui Chang, Dau-Ming Niu, Chia-Ying Chang, Pao-Chin Chiu, Yen-Yin Chou, Hui-Pin Hsia, Meng-Che Tsai, Mei-Chyn Chao, Li-Ping Tsai, Chia-Feng Yang, Pen-Hua Su, Yu-Wen Pan, **Chen-Hao Lee**, Tzu-Hung Chu, Chih-Kuang Chuang, Shuan-Pei Li. Epigenotype, Genotype, and Phenotype Analysis of Taiwanese Patients with Silver-Russell Syndrome. Journal of Personalized Medicine, 2021 Nov 13;11(11):1197.
Poster presentation	
1.	**Chen-Hao Lee**, Hsiang-Yu Lin, Chih-Kuang Chuang, Peih-Shan Wu, Chih-Ping Chen, Shuan-Pei Lin. Array Based Comparative Genomic Hybridization Applying for Multiple Congenital Anomalies, Developmental Delay /Intellectual Disability, Autism Spectrum Disorders, and Seizure Disorder in Taiwan. European Human Genetics Conference 2015, June 6 - 9, 2015 in Glasgow, Scotland, United Kingdom.
2.	**Chen-Hao Lee**, Yung-Ning Yang, Po-Jui Ko, Ming-Lun, Yeh, Li-Min Chen, Yu-Tsun Su, Ni-Chung Lee. A missense mutation of COL2A1 gene in an achondrogensis type II neonate (Case report). American Society of Human Genetics 2018 Annual Meeting, October 6-20, 2018 in Vancouver, Canada.
3.	**Chen-Hao Lee**, Li-Min Chen, Yu-Tsun Su, Yin-Hsiu Chien, Wuh-Liang Hwu, Ni-Chung Lee. Application of Exome Sequencing for Intellectual Disability and/or Multiple Congenital Anomalies. 2019 Asia Pacific Conference on Human Genetics, November 7-9, 2019 in Manila, Philippines.