### CONTENTS

**EDITORIAL**

8432  Evolution of *World Journal of Clinical Cases* over the past 5 years  
*Mathu S*

**OPINION REVIEW**

8436  NF-κB: A novel therapeutic pathway for gastroesophageal reflux disease?  
*Zhang ML, Ran LQ, Wu MJ, Jia QC, Qin ZM, Peng YG*

**MINIREVIEWS**

8443  Obligate aerobic, gram-positive, weak acid-fast, nonmotile bacilli, *Tsukamurella tyrosinosolvens*: Minireview of a rare opportunistic pathogen  

8450  Diffusion tensor imaging pipeline measures of cerebral white matter integrity: An overview of recent advances and prospects  
*Safri AA, Nassir CMNCM, Iman IN, Mohd Taib NH, Achuthan A, Mustapha M*

8463  Graft choices for anterolateral ligament knee reconstruction surgery: Current concepts  
*Chalidis B, Pitsilos C, Kitridis D, Givissis P*

8474  Overview of the anterolateral complex of the knee  
*Garcia-Mansilla I, Zicaro JP, Martinez EF, Astou J, Yacuzzi C, Costa-Paz M*

8482  Complication of lengthening and the role of post-operative care, physical and psychological rehabilitation among fibula hemimelia  
*Salimi M, Sarallah R, Javanshir S, Mirghaderi SP, Salimi A, Khanzadeh S*

**ORIGINAL ARTICLE**

**Clinical and Translational Research**

8490  Pyroptosis-related genes play a significant role in the prognosis of gastric cancer  
*Guan SH, Wang XY, Shang P, Du QC, Li MZ, Xing X, Yan B*

**Retrospective Study**

8506  Effects of propofol combined with lidocaine on hemodynamics, serum adrenocorticotropic hormone, interleukin-6, and cortisol in children  
*Shi S, Gan L, Jin CN, Liu RF*

8514  Correlation analysis of national elite Chinese male table tennis players’ shoulder proprioception and muscle strength  
*Shang XD, Zhang EM, Chen ZL, Zhang L, Qian JH*
8525 Clinical value of contrast-enhanced ultrasound in early diagnosis of small hepatocellular carcinoma (≤ 2 cm)

Mei Q, Yu M, Chen Q

8535 Identification of predictive factors for post-transarterial chemoembolization liver failure in hepatocellular carcinoma patients: A retrospective study

Yuan M, Chen TY, Chen XR, Lu YF, Shi J, Zhang WS, Ye C, Tang BZ, Yang ZG

8547 Clinical significance of half-hepatic blood flow occlusion technology in patients with hepatocellular carcinoma with cirrhosis

Liu D, Fang JM, Chen XQ

8556 Which octogenarian patients are at higher risk after cholecystectomy for symptomatic gallstone disease? A single center cohort study

D’Acapito F, Solaini L, Di Pietrantonio D, Taucer F, Mirarchi MT, Antiemi E, Flamini F, Amato A, Framarini M, Ercolani G

Clinical Trials Study

8568 Computed tomography combined with gastroscopy for assessment of pancreatic segmental portal hypertension

Wang YL, Zhang HW, Lin F

Observational Study

8578 Psychological needs of parents of children with complicated congenital heart disease after admitting to pediatric intensive care unit: A questionnaire study

Zhu JH, Jin CD, Tang XM

Prospective Study

8587 Quantitative differentiation of malignant and benign thyroid nodules with multi-parameter diffusion-weighted imaging


Randomized Controlled Trial

8599 Application of unified protocol as a transdiagnostic treatment for emotional disorders during COVID-19: An internet-delivered randomized controlled trial

Yan K, Yusufi MH, Nazari N

8615 High-flow nasal cannula oxygen therapy during anesthesia recovery for older orthopedic surgery patients: A prospective randomized controlled trial

Li XN, Zhou CC, Lin ZQ, Jia B, Li XY, Zhao GF, Ye F

SYSTEMATIC REVIEWS

8625 Assessment tools for differential diagnosis of neglect: Focusing on egocentric neglect and allocentric neglect

Lee SH, Lim BC, Jeong CY, Kim JH, Jang WH
CASE REPORT

8634 Exome analysis for Cronkhite-Canada syndrome: A case report
Li ZD, Rong L, He YJ, Ji YZ, Li X, Song FZ, Li XA

8641 Discrepancy between non-invasive prenatal testing result and fetal karyotype caused by rare confined placental mosaicism: A case report
Li Z, Lai GR

8648 Paroxysmal speech disorder as the initial symptom in a young adult with anti-N-methyl-D-aspartate receptor encephalitis: A case report
Hu CC, Pan XL, Zhang MX, Chen HF

8656 Anesthetics management of a renal angiomyolipoma using pulse pressure variation and non-invasive cardiac output monitoring: A case report
Jeon WJ, Shin WJ, Yoon YJ, Park CW, Shim JH, Cho SY

8662 Traumatic giant cell tumor of rib: A case report
Chen YS, Kao HW, Huang HY, Huang TW

8667 Analysis of two naval pilots' ejection injuries: Two case reports
Zeng J, Liu XP, Yi JC, Lu X, Liu DD, Jiang YQ, Liu YB, Tian JQ

8673 Beware of the DeBakey type I aortic dissection hidden by ischemic stroke: Two case reports
Chen SQ, Luo WL, Liu W, Wang LZ

8679 Unilateral lichen planus with Blaschko line distribution: A case report
Dong S, Zhu WJ, Xu M, Zhao XQ, Mou Y

8686 Clinical features and progress of ischemic gastritis with high fatalities: Seven case reports

8690 Retinoblastoma in an older child with secondary glaucoma as the first clinical presenting symptom: A case report
Zhang Y, Tang L

8703 Recurrent herpes zoster in a rheumatoid arthritis patient treated with tofacitinib: A case report and review of the literature
Lin QX, Meng JJ, Pang YY, Qu Y

8709 Intra-abdominal ectopic bronchogenic cyst with a mucinous neoplasm harboring a GNAS mutation: A case report

8718 Effects of intravascular photobiomodulation on motor deficits and brain perfusion images in intractable myasthenia gravis: A case report
Lan CH, Wu YC, Chiang CC, Chang ST
<table>
<thead>
<tr>
<th>Page</th>
<th>Title</th>
<th>Authors</th>
</tr>
</thead>
<tbody>
<tr>
<td>8728</td>
<td>Spontaneous acute epidural hematoma secondary to skull and dural metastasis of hepatocellular carcinoma: A case report</td>
<td>Lv GZ, Li GC, Tang WT, Zhou D, Yang Y</td>
</tr>
<tr>
<td>8735</td>
<td>Malignant melanotic nerve sheath tumors in the spinal canal of psammomatous and non-psammomatous type: Two case reports</td>
<td>Yeom JA, Song YS, Lee IS, Han IH, Choi KU</td>
</tr>
<tr>
<td>8742</td>
<td>When should endovascular gastrointestinal anastomosis transection Glissonean pedicle not be used in hepatectomy? A case report</td>
<td>Zhao J, Dang YL</td>
</tr>
<tr>
<td>8749</td>
<td>VARS2 gene mutation leading to overall developmental delay in a child with epilepsy: A case report</td>
<td>Wu XH, Lin SZ, Zhou YQ, Wang WQ, Li JY, Chen QD</td>
</tr>
<tr>
<td>8755</td>
<td>Junctional bradycardia in a patient with COVID-19: A case report</td>
<td>Aedh AI</td>
</tr>
<tr>
<td>8768</td>
<td>High scored thyroid storm after stomach cancer perforation: A case report</td>
<td>Baik SM, Pae Y, Lee JM</td>
</tr>
<tr>
<td>8775</td>
<td>Cholecystitis-an uncommon complication following thoracic duct embolization for chylothorax: A case report</td>
<td>Dung LV, Hien MM, Tra My TT, Lau DT, Linh LT, Duc NM</td>
</tr>
<tr>
<td>8782</td>
<td>Endometrial squamous cell carcinoma originating from the cervix: A case report</td>
<td>Shu XY, Dai Z, Zhang S, Yang HX, Bi H</td>
</tr>
<tr>
<td>8788</td>
<td>Type 2 autoimmune pancreatitis associated with severe ulcerative colitis: Three case reports</td>
<td>Ghali M, Bensted K, Williams DB, Ghaly S</td>
</tr>
</tbody>
</table>

**LETTER TO THE EDITOR**

<table>
<thead>
<tr>
<th>Page</th>
<th>Title</th>
<th>Authors</th>
</tr>
</thead>
<tbody>
<tr>
<td>8805</td>
<td>Comment on “Posterior reversible encephalopathy syndrome in a patient with metastatic breast cancer: A case report”</td>
<td>Kunić S, Ibrahimagić OĆ, Kojić B, Džananović D</td>
</tr>
</tbody>
</table>
## About Cover

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## Aims and Scope

The primary aim of *World Journal of Clinical Cases (WJCC, World J Clin Cases)* is to provide scholars and readers from various fields of clinical medicine with a platform to publish high-quality clinical research articles and communicate their research findings online.

*WJCC* mainly publishes articles reporting research results and findings obtained in the field of clinical medicine and covering a wide range of topics, including case control studies, retrospective cohort studies, retrospective studies, clinical trials studies, observational studies, prospective studies, randomized controlled trials, randomized clinical trials, systematic reviews, meta-analysis, and case reports.

## Indexing/Abstracting

The *WJCC* is now abstracted and indexed in Science Citation Index Expanded (SCIE, also known as SciSearch®), Journal Citation Reports/Science Edition, Current Contents®/Clinical Medicine, PubMed, PubMed Central, Scopus, Reference Citation Analysis, China National Knowledge Infrastructure, China Science and Technology Journal Database, and Superstar Journals Database. The 2022 Edition of Journal Citation Reports® cites the 2021 impact factor (IF) for *WJCC* as 1.534; IF without journal self cites: 1.491; 5-year IF: 1.599; Journal Citation Indicator: 0.28; Ranking: 135 among 172 journals in medicine, general and internal; and Quartile category: Q4. The *WJCC*’s CiteScore for 2021 is 1.2 and Scopus CiteScore rank 2021: General Medicine is 443/826.

## Responsible Editors for This Issue

Production Editor: Ying-Yi Yuan; Production Department Director: Xu Guo; Editorial Office Director: Jin-Lei Wang.
Exome analysis for Cronkhite-Canada syndrome: A case report

Zhao-Dong Li, Li Rong, Yuan-Jing He, Yu-Zhu Ji, Xiang Li, Fang-Zhou Song, Xiao-An Li

**Abstract**

**BACKGROUND**

Cronkhite-Canada syndrome (CCS) is a rare, non-genetic disorder characterized by multiple gastrointestinal polyps, and ectodermal lesions such as alopecia, fingernail atrophy, and skin mucosal pigmentation. Unfortunately, the pathogenesis of CCS is currently unknown.

**CASE SUMMARY**

Here, we describe the case of an elderly female with diarrhea, fatigue, and hair loss, who experienced abdominal pain for over half a year and was found to have multiple gastrointestinal polyps. She was diagnosed with CCS and was treated with albumin supplementation and prednisone, and her electrolyte imbalance was corrected. Following treatment, her symptoms significantly improved. To elucidate the role of potential genetic events in the pathogenesis of CCS, we performed exome sequencing using an extract of her colorectal adenoma.

**CONCLUSION**

Our data revealed multiple somatic mutations and copy number variations. Our findings provide a novel insight into the potential mechanisms of CCS etiology.
Key Words: Whole exome sequencing; Cronkhite-Canada syndrome; Somatic mutations; Case report

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Core Tip: Cronkhite-Canada syndrome (CCS) is a rare, non-genetic disorder characterized by multiple gastrointestinal polyps, ectodermal lesions including alopecia, fingernail atrophy, and skin mucosal pigmentation. However, its pathogenesis is unclear. We performed exome sequencing in an elderly female patient and obtained somatic mutations in the hope that these data could provide a genetic perspective on the pathogenesis of CCS.

INTRODUCTION

Cronkite and Canada reported the first case of Cronkhite-Canada syndrome (CCS) in 1955, which was characterized by multiple gastrointestinal polyps, ectodermal lesions including alopecia, fingernail atrophy, and skin mucosal pigmentation[1]. Unfortunately, despite much research, the pathogenesis of CCS remains unknown. Some studies have reported an association between CCS and immune factors, especially infiltration of IgG4-positive plasma cells[2]. To date, there is no standard treatment for CCS; however, some studies have recommended glucocorticoid therapy for partial symptom relief[3]. Herein, we describe the case of an elderly female with the diagnosis of CCS. As part of her treatment regimen, she received albumin supplementation and prednisone, and her electrolyte disturbance was corrected. After 15 days, her symptoms, namely, hypokalemia and diarrhea, significantly improved. To elucidate the role of potential genetic events in the pathogenesis of CCS, we performed exome sequencing using DNA extracted from a colorectal adenoma. Our data revealed multiple somatic mutations and copy number variations. We are the first to identify 3 novel genetic mutations (USP24, KCNQ5, and FKBP10) in a CCS patient. Moreover, we demonstrated that the HPSE2, SPATA7, and ZC3H18 genes had markedly elevated copy numbers. Given this evidence, we hypothesized that these specific gene mutations and copy number variations are associated with the pathogenesis of CCS.

CASE PRESENTATION

Chief complaints
An elderly female patient sought treatment for diarrhea, fatigue, and hair loss, as well as abdominal pain for half a year at the Department of Gastroenterology on October 11, 2019.

History of present illness
She reported diarrhea 4-5 times daily, watery stools, no blood, and abdominal pain under the xiphoid process, not related to eating. Physical examination revealed obvious emaciation, alopecia, and nail atrophy.

History of past illness
Her past medical history is unremarkable.

Personal and family history
Her family history is unremarkable.

Physical examination
Physical examination revealed obvious emaciation, alopecia, and nail atrophy (Figure 1).

Laboratory examinations
Laboratory tests revealed the following: (routine blood examination) white blood cells: $10.03 \times 10^9$/L, hemoglobin: 95 g/L, and platelets: $151 \times 10^9$/L; (liver function) albumin 25 g/L; (electrolytes) Na+: 128 mmol/L, K+: 2.8 mmol/L.
Li ZD et al. Exome analysis for Cronkhite-Canada syndrome

Figure 1 Signs in the patient with Cronkhite-Canada syndrome. A: The patient had typical symptoms of hair loss (orange arrow); B: The patient had typical nail atrophy (orange arrow).

Imaging examinations
She underwent gastrointestinaloscopy, which revealed multiple small polyps in the gastrointestinal tract (Figure 2).

Further diagnostic work-up
DNA samples: This study was performed in accordance with the guidelines of the Ethics Committee of the Mianyang Central Hospital. DNA was extracted from a sample of the patient's colonic adenomas and normal colon tissue. The samples were obtained via endoscopic mucosal resection and were preserved in liquid nitrogen.

Whole exome sequencing: Genomic DNA ≥ 1.5 μg from colonic adenomas and normal colon tissue was used for whole exome sequencing library construction. The Agilent liquid chip capture system was employed for the efficient enrichment of human DNA in all exon regions. Upon library construction, Qubit2.0 was used for preliminary quantification, and the library was diluted to 1 ng/μL. Subsequently, Agilent 2100 was utilized for detection of the library insert size to ensure library quality (the effective concentration of the library was > 2 nmol/L). Next, PE150 high-throughput sequencing was performed based on the Illumina Hiseq platform. Finally, the library and capture experiment were constructed using the Agilent SureSelect v5 kit, according to the manufacturer's instructions.

Quality control: Upon collection of the original sequenced reads (Sequenced Reads), Cutadapt software was used to remove the adapter and reads with N (N denotes that the base information cannot be determined) ratio ≥ 10%. In addition, the read pairs were discarded if the low-quality bases (quality score ≤ 5) accounted for 50% of the entire single read. Furthermore, quality controls (including, sequencing error rate distribution, base quality distribution, and GC content distribution) were performed, based on the selected filtered reads using the above methods.

Data analysis: The BWA (v0.7.15) software was used to map the sequenced reads to the human reference genome GRCh37 (hg19). Picard software was used to remove the sequence generated by PCR-duplication. The somatic single nucleotide variations (SNVs) and InDel detection were performed using the GATK (v4.1.0.0) and muTect2 software, respectively. The ANNOVAR software (Mon, 17 Jul 2017) was used to annotate gene mutations. Lastly, the control-free software (v11.4) was used for copy number variation analysis.

Exon sequencing results
Somatic mutation analysis, based on samples from the CCS patient, identified 47 SNVs. 75% of them were nonsynonymous SNVs. In addition, approximately 70% of them occurred in exonic regions (Figure 3). Mutations of the USP24, KCNQ5, and FKBP10 genes were identified as deleterious via SIFT, LRT, Polyphen2, and Mutation Taster software (Supplementary Table 1). Based on the COSMIC data, the three mutations in genes USP24, KCNQ5, and FKBP10 may be novel in CCS. The mutated genes are associated with the regulation of DNA-templated transcription (Figure 4). Our analysis showed that the HPSE2, SPATA7, and ZC3H18 genes had markedly elevated copy numbers, while other significant altered fragments were located in the intergenic regions (Table 1).
Table 1 The identified copy number alterations, as shown by exome sequencing

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<th>Start</th>
<th>End</th>
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</table>

Figure 2 Endoscopic characteristics of the patient with Cronkhite-Canada syndrome. A: The patient presented with multiple adenomas of the stomach greater curvature (blue arrow); B: The patient presented with multiple adenomas of the colon (blue arrow).

**FINAL DIAGNOSIS**

The patient was diagnosed with CCS.

**TREATMENT**

As part of her treatment regimen, she received albumin supplementation 10 g/d and prednisone 10 mg/d, and her electrolyte imbalance was corrected.
OUTCOME AND FOLLOW-UP

Treatment lasted for 15 d and her diarrhea, hypokalemia, and abdominal pain significantly improved.

DISCUSSION

Approximately 500 cases of CCS are currently described in the literature, with an estimated incidence of 1 in 1 million per year. The average age at CCS diagnosis is early 60s and it is predominantly diagnosed in males[4]. At present, there is no definite treatment plan for CCS. Previous literature reported that glucocorticoids and immunosuppression have certain benefits, and some studies reported that anti-tumor necrosis factor therapy is effective in CCS patients[5]. Our treatment plan and effect are consistent with these published works. However, there are limited studies on the pathogenesis of CCS. Therefore, based on the fact that CCS is an acquired non-genetic disease, we performed exon sequencing of excised diseased and non-diseased tissues from our CCS patient, in an attempt to elucidate CCS pathogenesis. Based on our exon sequencing results, three genes, namely, USP24, KCNQ5, and FKBP10 may be related to CCS pathogenesis.

We searched Human Gene Mutation Database (HGMD) and found that USP24 belongs to a large family of cysteine proteinases that function as deubiquitinating enzymes. USP24 stabilizes the bromine domain protein and promotes malignant lung cancer[6]. In addition, Zhang L et al[7] found that USP24 deubiquitinasate regulates DNA damage by directly targeting the tumor suppressor gene p53. Also, the USP24-Mcl-1 axis may represent a novel strategy in treating acute T cell lymphoma[8], whereas functional studies of the FKBP10 mutation reported an association with osteogenesis imperfecta[9]. The above two genes are both related to tumor formation and body development. Combined with the clinicopathological characteristics of CCS, we speculated that gene mutations are involved in the formation of multiple intestinal adenomas. The KCNQ family protein activates slowly during depolarization and forms heterogeneous channels with the protein encoded by KCNQ5 gene. KCN5 dependent potassium channels play an important role in airway smooth muscle relaxation[10]. Given the symptoms of diarrhea and difficult-to-correct hypokalemia of CCS, the KNQ3 mutation seems to suggest an association.

In terms of copy variation, ZC3H18 copy number losses are known to contribute to homologous recombination defects in high-grade serous ovarian cancers[11]. HPSE2 was reported to play an inhibitory role in bladder cancer[12]. Mutations in SPATA7 are associated with fundus macular degeneration[13]. CCS patients have multiple clinicopathological manifestations, such as, multiple gastrointestinal adenomas, nail atrophy, skin pigmentation, and alopecia, which may be related to the increased copy number of the three genes mentioned above.

However, due to the isolation of individual cases, the sample size of phenotypic alterations, caused by the above gene mutations, needs to be further expanded.

CCS is a rare disease and its etiology is unclear. The autoimmune etiology of CCS was previously proposed, and case reports described beneficial responses to immunosuppressive therapies such as azathioprine, anti-tumor necrosis factor antibodies, cyclosporine, and sirolimus[14,15]. Interestingly, Brigid S. Boland also conducted exome sequencing on tissue from a CCS patient who responded effectively to infliximab, and found that PRKDC mutations may be involved. However, this gene was not included in our analysis[14]. This suggests that the data on these mutations require further validation using tissues from a large CCS patient population.
In conclusion, we report a classic case of CCS, which was effectively treated with parenteral nutritional support and glucocorticoids, and we explored the pathogenesis of CCS from the perspective of gene mutation. Based on our analysis, we identified several gene mutations and alterations in gene copy numbers. However, we acknowledge that more genetic and epidemiological research is necessary to understand the complex pathogenesis of this rare but highly fatal disease.

FOOTNOTES

Author contributions: Li ZD and He YJ treated the patient and drafted the manuscript; Rong L and Ji YZ performed the exon sequencing work; Li X helped to search for references; Zhou SF and Li XA performed data retrieval and helped with drafting of the manuscript; all authors read and approved the final manuscript.

Informed consent statement: Informed consent was obtained from the patient’s family for publication of this case.
Li ZD et al. Exome analysis for Cronkhite-Canada syndrome

Conflict-of-interest statement: The authors declare no conflicts of interest.


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REFERENCES
