OPINION REVIEW

7620 Whipple’s operation with a modified centralization concept: A model in low-volume Caribbean centers
Cawich SO, Pearce NW, Naraynsingh V, Shakla P, Deshpande RR

REVIEW

7631 Role of micronutrients in Alzheimer’s disease: Review of available evidence
Fei HX, Qian CF, Wu XM, Wei YH, Huang JY, Wei LH

MINIREVIEWS

7642 Application of imaging techniques in pancreaticobiliary maljunction
Wang JY, Mu PY, Xu YK, Bai YY, Shen DH

7653 Update on gut microbiota in gastrointestinal diseases
Nishida A, Nishino K, Ohno M, Sakai K, Owaki Y, Noda Y, Imaeda H

7665 Vascular complications of pancreatitis
Kalas MA, Leon M, Chavez LO, Canalizo E, Sarani S

ORIGINAL ARTICLE

Clinical and Translational Research

7674 Network pharmacology and molecular docking reveal zedoary turmeric-trisomes in Inflammatory bowel disease with intestinal fibrosis
Zheng L, Ji YY, Dai YC, Wen XL, Wu SC

Case Control Study

7686 Comprehensive proteomic signature and identification of CDKN2A as a promising prognostic biomarker and therapeutic target of colorectal cancer
Wang QQ, Zhou YC, Zhou Ge YJ, Qin G, Yin TF, Zhao DY, Tan C, Yao SK

Retrospective Cohort Study

7698 Is anoplasty superior to scar revision surgery for post-hemorrhoidectomy anal stenosis? Six years of experience
Wang YT, Chu KJ, Lin KH, Chang CK, Kang JC, Chen CY, Hu JM, Pu TW

Retrospective Study

7708 Short- (30-90 days) and mid-term (1-3 years) outcomes and prognostic factors of patients with esophageal cancer undergoing surgical treatments
Shi MK, Mei YQ, Shi JI
<table>
<thead>
<tr>
<th>Page</th>
<th>Title</th>
<th>Authors</th>
</tr>
</thead>
<tbody>
<tr>
<td>7720</td>
<td>Effectiveness of pulsed radiofrequency on the medial cervical branches for cervical facet joint pain</td>
<td>Chang MC, Yang S</td>
</tr>
<tr>
<td>7738</td>
<td>Correlation between the warning symptoms and prognosis of cardiac arrest</td>
<td>Zheng K, Bai Y, Zhai QR, Du LF, Ge HX, Wang GX, Ma QB</td>
</tr>
<tr>
<td>7749</td>
<td>Serum ferritin levels in children with attention deficit hyperactivity disorder and tic disorder</td>
<td>Tang CY, Wen F</td>
</tr>
<tr>
<td>7760</td>
<td>Application of metagenomic next-generation sequencing in the diagnosis of infectious diseases of the central nervous system after empirical treatment</td>
<td>Chen YY, Guo Y, Xue XH, Pang F</td>
</tr>
<tr>
<td>7785</td>
<td>Prospective single-center feasible study of innovative autorelease bile duct supporter to delay adverse events after endoscopic papillectomy</td>
<td>Liu SZ, Chai NL, Li HK, Feng XX, Zhai YQ, Wang NJ, Gao Y, Gao F, Wang SS, Linghu EQ</td>
</tr>
</tbody>
</table>

**Clinical Trials Study**

<table>
<thead>
<tr>
<th>Page</th>
<th>Title</th>
<th>Authors</th>
</tr>
</thead>
<tbody>
<tr>
<td>7794</td>
<td>Performance of Dexcom G5 and FreeStyle Libre sensors tested simultaneously in people with type 1 or 2 diabetes and advanced chronic kidney disease</td>
<td>Ölafsdóttir AF, Andelin M, Saeed A, Sofizadeh S, Hamoodi H, Jansson PA, Lind M</td>
</tr>
</tbody>
</table>

**Observational Study**

<table>
<thead>
<tr>
<th>Page</th>
<th>Title</th>
<th>Authors</th>
</tr>
</thead>
<tbody>
<tr>
<td>7808</td>
<td>Complications of chronic pancreatitis prior to and following surgical treatment: A proposal for classification</td>
<td>Murruste M, Kirsimägi Ü, Kase K, Veršinina T, Talving P, Lepner U</td>
</tr>
<tr>
<td>7825</td>
<td>Effects of comprehensive nursing on postoperative complications, mental status and quality of life in patients with glioma</td>
<td>Dong H, Zhang XL, Deng CX, Luo B</td>
</tr>
</tbody>
</table>

**Prospective Study**

<table>
<thead>
<tr>
<th>Page</th>
<th>Title</th>
<th>Authors</th>
</tr>
</thead>
<tbody>
<tr>
<td>7832</td>
<td>Predictors of long-term anxiety and depression in discharged COVID-19 patients: A follow-up study</td>
<td>Boyraz RK, Şahan E, Boylu ME, Korponar I</td>
</tr>
</tbody>
</table>

**META-ANALYSIS**

<table>
<thead>
<tr>
<th>Page</th>
<th>Title</th>
<th>Authors</th>
</tr>
</thead>
<tbody>
<tr>
<td>Page</td>
<td>Title</td>
<td>Authors</td>
</tr>
<tr>
<td>------</td>
<td>----------------------------------------------------------------------</td>
<td>-------------------------------------------------------------------------</td>
</tr>
<tr>
<td>7859</td>
<td>Rectal nonsteroidal anti-inflammatory drugs, glyceryl trinitrate, or combinations for prophylaxis of post-endoscopic retrograde cholangiopancreatography pancreatitis: A network meta-analysis</td>
<td>Shi QQ, Huang GX, Li W, Yang JR, Ning XY</td>
</tr>
<tr>
<td>7872</td>
<td>Effect of celecoxib on improving depression: A systematic review and meta-analysis</td>
<td>Wang Z, Wu Q, Wang Q</td>
</tr>
<tr>
<td>7883</td>
<td>Rectal mature teratoma: A case report</td>
<td>Liu JL, Sun PL</td>
</tr>
<tr>
<td>7890</td>
<td>Antibiotic and glucocorticoid-induced recapitulated hematological remission in acute myeloid leukemia: A case report and review of literature</td>
<td>Sun XY, Yang XD, Yang XQ, Ju B, Xie NN, Xu J, Zhao XC</td>
</tr>
<tr>
<td>7899</td>
<td>Non-secretory multiple myeloma expressed as multiple extramedullary plasmacytoma with an endobronchial lesion mimicking metastatic cancer: A case report</td>
<td>Lee SB, Park CY, Lee HJ, Hong R, Kim WS, Park SG</td>
</tr>
<tr>
<td>7906</td>
<td>Latamoxef-induced severe thrombocytopenia during the treatment of pulmonary infection: A case report</td>
<td>Zhang RY, Zhang JJ, Li JM, Xu YY, Xu YH, Cai XJ</td>
</tr>
<tr>
<td>7913</td>
<td>Multicentric reticulohistiocytosis with prominent skin lesions and arthritis: A case report</td>
<td>Xu XL, Liang XH, Liu J, Deng X, Zhang L, Wang ZG</td>
</tr>
<tr>
<td>7924</td>
<td>Brainstem abscesses caused by Listeria monocytogenes: A case report</td>
<td>Wang J, Li YC, Yang KY, Wang J, Dong Z</td>
</tr>
<tr>
<td>7931</td>
<td>Primary hypertension in a postoperative paraganglioma patient: A case report</td>
<td>Wei JH, Yan HL</td>
</tr>
<tr>
<td>7936</td>
<td>Long-term survival of gastric mixed neuroendocrine-non-neuroendocrine neoplasm: Two case reports</td>
<td>Woo LT, Ding YF, Mao CY, Qian J, Zhang XM, Xu N</td>
</tr>
<tr>
<td>7944</td>
<td>Percutaneous transfemoral endoscopic decompression combined with percutaneous vertebroplasty in treatment of lumbar vertebral body metastases: A case report</td>
<td>Ran Q, Li T, Kuang ZP, Guo XH</td>
</tr>
<tr>
<td>7950</td>
<td>Atypical imaging features of the primary spinal cord glioblastoma: A case report</td>
<td>Liang XY, Chen YP, Li Q, Zhou ZW</td>
</tr>
<tr>
<td>7960</td>
<td>Resection with limb salvage in an Asian male adolescent with Ewing’s sarcoma: A case report</td>
<td>Lai CY, Chen KJ, Ho TY, Li LY, Kuo CC, Chen HT, Fong YC</td>
</tr>
</tbody>
</table>
## Contents

### Thrice Monthly Volume 10 Number 22 August 6, 2022

<table>
<thead>
<tr>
<th>Page</th>
<th>Title</th>
<th>Authors</th>
</tr>
</thead>
<tbody>
<tr>
<td>7973</td>
<td>Delayed arterial symptomatic epidural hematoma on the 14th day after posterior lumbar interbody fusion: A case report</td>
<td>Hao SS, Gao ZF, Li HK, Liu S, Dong SL, Chen HL, Zhang ZF</td>
</tr>
<tr>
<td>7982</td>
<td>Clinical and genetic analysis of nonketotic hyperglycinemia: A case report</td>
<td>Ning JJ, Li F, Li SQ</td>
</tr>
<tr>
<td>7994</td>
<td>Occurrence of MYD88L265P and CD79B mutations in diffuse large B cell lymphoma with bone marrow infiltration: A case report</td>
<td>Huang WY, Weng ZY</td>
</tr>
<tr>
<td>8003</td>
<td>Rare case of compartment syndrome provoked by inhalation of polyurethane agent: A case report</td>
<td>Choi JH, Oh HM, Hwang JH, Kim KS, Lee SY</td>
</tr>
<tr>
<td>8009</td>
<td>Acute ischemic Stroke combined with Stanford type A aortic dissection: A case report and literature review</td>
<td>He ZY, Yao LP, Wang XK, Chen NY, Zhao JJ, Zhou Q, Yang XF</td>
</tr>
<tr>
<td>8018</td>
<td>Compound-honeysuckle-induced drug eruption with special manifestations: A case report</td>
<td>Zhou LF, Lu R</td>
</tr>
<tr>
<td>8025</td>
<td>Spontaneous internal carotid artery pseudoaneurysm complicated with ischemic stroke in a young man: A case report and review of literature</td>
<td>Zhong YL, Feng JP, Luo H, Gong XH, Wei ZH</td>
</tr>
<tr>
<td>8034</td>
<td>Microcystic adnexal carcinoma misdiagnosed as a “recurrent epidermal cyst”: A case report</td>
<td>Yang SX, Mou Y, Wang S, Hu X, Li FQ</td>
</tr>
<tr>
<td>8040</td>
<td>Accidental discovery of appendiceal carcinoma during gynecological surgery: A case report</td>
<td>Wang L, Dong Y, Chen YH, Wang YN, Sun L</td>
</tr>
<tr>
<td>8045</td>
<td>Intra-ampullary papillary-tubular neoplasm combined with ampullary neuroendocrine carcinoma: A case report</td>
<td>Zavrtanik H, Lucas B, Tomažič A</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>8054</td>
<td>Commentary on &quot;Primary orbital monophasic synovial sarcoma with calcification: A case report&quot;</td>
<td>Tokar O, Aydin S, Karavas E</td>
</tr>
</tbody>
</table>
ABOUT COVER
Editorial Board Member of *World Journal of Clinical Cases*, Bennete Aloysius Fernandes, MDS, Professor, Faculty of Dentistry, SEGi University, Kota Damansara 47810, Selangor, Malaysia. drben17@yahoo.com

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: Xu Guo; Production Department Director: Xiang Li; Editorial Office Director: Jin-Lei Wang.

NAME OF JOURNAL
*World Journal of Clinical Cases*

ISSN
ISSN 2307-8960 (online)

LAUNCH DATE
April 16, 2013

FREQUENCY
Thrice Monthly

EDITORS-IN-CHIEF
Bao-Gan Peng, Jerzy Tadeusz Chudek, George Kontogeorgos, Maurizio Serati, Ja Hyeon Ku

EDITORIAL BOARD MEMBERS
https://www.wjgnet.com/2307-8960/editorialboard.htm

PUBLICATION DATE
August 6, 2022

COPYRIGHT
© 2022 Baishideng Publishing Group Inc

INSTRUCTIONS TO AUTHORS
https://www.wjgnet.com/bpg/gerinfo/204

GUIDELINES FOR ETHICS DOCUMENTS
https://www.wjgnet.com/bpg/gerInfo/287

GUIDELINES FOR NON-NATIVE SPEAKERS OF ENGLISH
https://www.wjgnet.com/bpg/gerInfo/240

PUBLICATION ETHICS
https://www.wjgnet.com/bpg/gerInfo/288

PUBLICATION MISCONDUCT
https://www.wjgnet.com/bpg/gerInfo/208

ARTICLE PROCESSING CHARGE
https://www.wjgnet.com/bpg/gerInfo/242

STEPS FOR SUBMITTING MANUSCRIPTS
https://www.wjgnet.com/bpg/gerInfo/239

ONLINE SUBMISSION
https://www.f6publishing.com
CASE REPORT

Latamoxef-induced severe thrombocytopenia during the treatment of pulmonary infection: A case report

Ruo-Ying Zhang, Jun-Jie Zhang, Jin-Meng Li, Ying-Ying Xu, Yue-Huan Xu, Xin-Jun Cai

Specialty type: Medicine, research and experimental

Provenance and peer review: Unsolicited article; Externally peer reviewed.

Peer-review model: Single blind

Peer-review report's scientific quality classification
Grade A (Excellent): 0
Grade B (Very good): 0
Grade C (Good): C
Grade D (Fair): D
Grade E (Poor): 0

P-Reviewer: Gaman MA, Romania; Socea B, Romania

Received: October 13, 2021
Peer-review started: October 13, 2021
First decision: January 11, 2022
Revised: January 24, 2022
Accepted: June 24, 2022
Article in press: June 24, 2022
Published online: August 6, 2022

Abstract

BACKGROUND
Latamoxef shows excellent antibacterial activity against anaerobic bacteria such as Bacteroides fragilis. Reports of thrombocytopenic toxicity of latamoxef are limited. This report presents a case of severe thrombocytopenia possibly induced by latamoxef, an infrequent adverse drug reaction in a young patient with tuberculosis and Crohn's disease in China.

CASE SUMMARY
We reported a case of severe thrombocytopenia induced by latamoxef in a 28-year-old man with tuberculosis and Crohn's disease. On admission, the patient presented with a cough productive of bloody sputum, a chest computed tomogram suggested scattered mottled, high-density shadows in both lungs. Laboratory tests indicated a platelet count of 140000/μL. Considered a pulmonary bacterial infection, the patient received anti-infection therapy with latamoxef (dose: 2.0 g) intravenously Q12h. On the 9th day of treatment, the platelet count decreased to 44000/μL. On the 12th day, scattered purpura and ecchymosis appeared on the patient’s limbs and trunk, and the platelet count decreased to 9000/μL after latamoxef treatment for 15 d. Three days after discontinuation of latamoxef, the platelet count recovered to 157000/μL, and the area of scattered purpura and ecchymosis on the limbs and trunk decreased. The platelet counts remained in the normal range, and no thrombocytopenia was found at follow-up 15 mo after discharge.

Ruo-Ying Zhang, Jin-Meng Li, Ying-Ying Xu, Xin-Jun Cai, Department of Pharmacy, Affiliated Hangzhou Chest Hospital, Zhejiang University School of Medicine, Hangzhou 310000, Zhejiang Province, China

Jun-Jie Zhang, Department of Out-patient, Zhejiang Medical and Health Group Hangzhou Hospital, Hangzhou 310000, Zhejiang Province, China

Yue-Huan Xu, Tuberculosis Treatment Centre, Affiliated Hangzhou Chest Hospital, Zhejiang University School of Medicine, Hangzhou 310000, Zhejiang Province, China

Corresponding author: Xin-Jun Cai, Doctor, Chief Pharmacist, Department of Pharmacy, Affiliated Hangzhou Chest Hospital, Zhejiang University School of Medicine, No. 208 East Huancheng Road, Xiaocheng District, Hangzhou 310000, Zhejiang Province, China.
zjtcmlxj@zcmu.edu.cn
CONCLUSION
For patients treated with latamoxef, platelet counts should be carefully followed, and caregivers should be vigilant for the appearance of scattered ecchymosis.

Key Words: Thrombocytopenia; Latamoxef; Adverse drug reactions; Young onset; Case report

Core Tip: We described a case of severe thrombocytopenia likely induced by latamoxef, an infrequent adverse drug reaction in a young patient with tuberculosis and Crohn's disease. We followed the changes in platelet counts and the appearance of purpura during latamoxef treatment and after drug withdrawal and excluded other possible causes of thrombocytopenia. Our findings suggested that the patient's thrombocytopenia was caused by latamoxef. This is the first reported case of severe thrombocytopenia induced by latamoxef in a young Chinese patient to the best of our knowledge.

Citation: Zhang RY, Zhang JJ, Li JM, Xu YY, Xu YH, Cai XJ. Latamoxef-induced severe thrombocytopenia during the treatment of pulmonary infection: A case report. World J Clin Cases 2022; 10(22): 7906-7912
URL: https://www.wjgnet.com/2307-8960/full/v10/i22/7906.htm
DOI: https://dx.doi.org/10.12998/wjcc.v10.i22.7906

INTRODUCTION
Latamoxef is a new semisynthetic oxacephem antibiotic structurally similar to third-generation cephalosporins. Latamoxef has excellent antibacterial activity against anaerobic bacteria such as Bacteroides fragilis. It is stable against β-lactamases produced by most Enterobacteriaceae, mediated by plasmids or partially by chromosomes[1]. The primary associated adverse reactions are rash, drug fever, hepatic and renal dysfunction, neutropenia, and eosinophilia, followed by coagulation dysfunction, with an incidence of 12.45%[2,3]. Thrombocytopenia is a common blood disorder characterized by the destruction of circulating platelets and inhibition of platelet production[4]. Although several studies have reported that latamoxef could cause thrombocytopenia[5-7], thrombocytopenia induced by latamoxef in the Chinese population is rare and clinicians often overlook latamoxef-induced thrombocytopenia. This case report presents the first case of severe thrombocytopenia and multiple ecchymoses caused by latamoxef in a young Chinese patient. We defined thrombocytopenia as a platelet count less than 100000/μL[8].

CASE PRESENTATION

Chief complaints
A 28-year-old male patient presented to the hospital with a fever for one month and a cough for more than ten days.

History of present illness
He developed a fever at about 38°C without obvious inducement one month prior and went to another hospital’s emergency department. He received a diagnosis of upper respiratory tract infection. The symptoms subsided after symptomatic treatment. More than 10 days before presentation, the patient had a paroxysmal cough with white sticky sputum and was diagnosed with pneumonia. Symptoms did not improve after expectorant treatment. In the days before the presentation, he had developed yellow and bloody sputum accompanied by night sweats.

History of past illness
The patient had a history of Crohn’s disease for more than five years and took mesalazine sustained-release tablets. Half a year prior, he stopped the mesalazine and switched to adalimumab injection once every two weeks, and he was in stable condition at presentation.

Personal and family history
There is no specific family history of illness.
Physical examination
Several enlarged lymph nodes were found on the left and right sides of the patient's neck. The skin color was normal without ecchymosis, and respiratory rate and vital signs were normal.

Laboratory examinations
White blood cell (WBC) count was below the normal range, while hemoglobin (HGB) and platelet count were at normal levels (Figure 1). Other test indicators were in the normal ranges.

Imaging examinations
A chest computed tomogram suggested scattered mottled, high-density shadows in both lungs, mediastinal and hilar lymph node enlargement, and several nodules in the spleen.

FINAL DIAGNOSIS
Secondary tuberculosis with sputum smear-negative, initial treatment; Cervical lymphatic tuberculosis; Splenic tuberculosis; Crohn's disease; Thrombocytopenia; Leukopenia.

TREATMENT
Because the diagnosis of tuberculosis was not clear initially, we considered it a bacterial infection. The patient first received anti-infective therapy with latamoxef (2.0 g) intravenously every 12 h and leucogen tablets (20.0 mg) three times per day for leukocytopenia. The timeline of the overall treatment process is presented in Table 1.

On day 9 (18:00 h) after initiation of latamoxef treatment, the patient developed chills and fever to 38.2 °C without shivering and cough with a small amount of sputum. On day 10, after initiation of latamoxef treatment, the patient received isoniazid tablets (0.3 g/d) and rifampicin capsules 0.6 g daily, considering his history of immunosuppressive agents and positive T SPOT-TB testing results, and latent infection with Mycobacterium tuberculosis was evident. On day 11, cervical lymph node aspirate fluid grew Mycobacterium tuberculosis complex sensitive to rifampicin. Pathological examination of a biopsy specimen from a left cervical lymph node revealed chronic granulomatous lymphadenitis with coagulative necrosis. Considering the presence of secondary pulmonary tuberculosis, cervical lymph node tuberculosis, and splenic tuberculosis, we added pyrazinamide 0.5 g three times per day and ethambutol 1.0 g daily in combination with isoniazid and rifampicin.

On the 12th day, the patient’s body temperature returned to normal but scattered purpura and ecchymosis appeared on his limbs and trunk. The platelet count decreased to 7000/μL. Considering that this might be thrombocytopenia induced by rifampicin, we replaced rifampicin with levofloxacin sodium chloride injection, 0.5 g intravenous drip once a day. Following consultation with hematology, we added subcutaneous injection of recombinant human thrombopoietin at 15000 units per day and intravenous infusion of human immunoglobulin (20.0 g/d), 15 units of platelets, and 5 mg of dexamethasone. The patient developed hemoptysis on day 13, and we added intravenous infusion of tranexamic acid sodium chloride (0.5 g/d), etamsylate (2.0 g/d) and spearhead agkistrodon hemocogulase (2.0 U/d) for hemostasis.

On the 15th day, the platelet count decreased to 9000/μL, suggesting that the patient was in a critical state. Because the patient could not afford the medications, pharmacists simplified the prescriptions. We recommended discontinuing latamoxef 2.0 g Q12H and adding an intravenous injection of 10 mg of vitamin K1 once a day, and the clinicians agreed. On the 16th day, the platelet count increased to 57000/μL. We discontinued the human immunoglobulin injection and recombinant human thrombopoietin. On the 17th day, the platelet count rapidly recovered to 157000/μL. We discontinued vitamin K1 and dexamethasone. Since then, the patient did not use latamoxef and was discharged on the 24th day taking isoniazid, ethambutol, pyrazinamide, and levofloxacin for tuberculosis treatment. Figure 1 display the fluctuation of peripheral blood WBC, HGB, and platelets, respectively, along with medications. Figure 2 display the ecchymoses before discontinuation of latamoxef and before discharge, respectively.

OUTCOME AND FOLLOW-UP
The patient was followed up at the first, third, and fifth week and monthly after discharge. The platelet counts and the HGB concentrations remained stable and in the normal range. Prothrombin and activated partial thromboplastin were normal from admission to platelet recovery. No thrombocytopenia was found at follow-up 15 mo after discharge.
Table 1 Timeline of the treatment process

<table>
<thead>
<tr>
<th>Time</th>
<th>Symptom</th>
<th>Platelet counts</th>
<th>Treatment</th>
</tr>
</thead>
<tbody>
<tr>
<td>Day 1</td>
<td>Bloody sputum, scattered mottled, high-density shadows in both lungs</td>
<td>140000/μL</td>
<td>Latamoxef (dose: 2.0 g) intravenously Q12H</td>
</tr>
<tr>
<td>Day 9</td>
<td>Chills and fever to 38.2 °C (18:00)</td>
<td>44000/μL (8:00 am)</td>
<td>-</td>
</tr>
<tr>
<td>Day 10</td>
<td>Positive T SPOT-TB testing results</td>
<td>-</td>
<td>Added isoniazid tablets 0.3 g QD, rifampicin capsules 0.6 g QD</td>
</tr>
<tr>
<td>Day 11</td>
<td>Secondary pulmonary tuberculosis, cervical lymph node tuberculosis, and splenic tuberculosis were confirmed</td>
<td>-</td>
<td>Continued adding pyrazinamide 0.5 g TID, ethambutol 1.0 g QD</td>
</tr>
<tr>
<td>Day 12</td>
<td>Body temperature returned to normal but scattered purpura and ecchymosis appeared on his limbs and trunk’s skin</td>
<td>7000/μL</td>
<td>Replaced rifampicin with levofloxacin; Added recombinant human thrombopoietin (15000 units/d), human immunoglobulin (20.0 g/d), 15 units platelets, and 5 mg dexamethasone</td>
</tr>
<tr>
<td>Day 13</td>
<td>Hemoptysis</td>
<td>44000/μL</td>
<td>Continued adding tranexamic acid sodium chloride (0.5 g/d), etamsylate (2.0 g/d) and spearhead agkistrodon hemocoagulase (2.0 U/d) for hemostasis</td>
</tr>
<tr>
<td>Day 15</td>
<td>Critical state</td>
<td>9000/μL</td>
<td>Discontinued latamoxef 2.0 g Q12H and added vitamin K1 (10 mg/d)</td>
</tr>
<tr>
<td>Day 16</td>
<td>-</td>
<td>57000/μL</td>
<td>Discontinued the human immunoglobulin injection and recombinant human thrombopoietin</td>
</tr>
<tr>
<td>Day 17</td>
<td>-</td>
<td>157000/μL</td>
<td>Discontinued vitamin K1 and dexamethasone</td>
</tr>
<tr>
<td>Day 23</td>
<td>-</td>
<td>255000/μL</td>
<td>-</td>
</tr>
<tr>
<td>Day 24</td>
<td>Discharged</td>
<td>-</td>
<td>Took isoniazid, ethambutol, pyrazinamide, and levofloxacin for tuberculosis treatment</td>
</tr>
<tr>
<td>The 1, 3, 5 wk, and 15-mo after discharge</td>
<td>-</td>
<td>Normal</td>
<td>Took isoniazid, ethambutol, pyrazinamide, and levofloxacin for tuberculosis treatment</td>
</tr>
</tbody>
</table>

Thrombocytopenia: Platelet count less than 100000/μL. Abnormal values are given in italic font.

DISCUSSION

Our patient’s thrombocytopenia induced by latamoxef was unique. To our best knowledge, this is the first documented case in a young Chinese patient. Vayne et al[9] reported that drug-mediated immune thrombocytopenia often gave rise to a higher risk of bleeding. Generally, thrombocytopenia occurs after 5 to 10 d of drug exposure, and the median platelet count is usually less than 20000/μL. Platelet counts usually begin to recover at four to five half-lives or within two to three days after discontinuation[8,9]. The literature suggested that rifampicin had a strong tendency to cause thrombocytopenia with an incidence of between 1% and 10%[10]. A systematic evaluation of 153 drugs conducted by Arnold et al[11] found that the most drugs contributing to drug-induced immune thrombocytopenia were rifampicin, quinine, vancomycin, and ceftriaxone.

The patient started oral rifampicin on the 10th day and stopped on the 12th day. We excluded rifampicin-induced immune thrombocytopenia based on the following criteria: (1) The time of occurrence was not in line with expectations. Before taking rifampicin, the patient received latamoxef alone. At that time, the platelet count decreased significantly from 140000/μL to 44000/μL (by 68.57%); (2) The exposure time of rifampicin was short (only two days), far less than the exposure time of five to ten days; this exposure was not sufficient to cause a decline in the platelet count[8,9,12]; and (3) The elimination half-life of rifampicin is three to five hours, and the patient had been off rifampicin for three days before the recurrence of thrombocytopenia; this time-course was inconsistent with the reported recovery of platelet counts after four to five half-lives. We transfused 15 units of platelets and administered human immunoglobulin, glucocorticoid after discontinuation of rifampicin to retard platelet clearance; however, the patient’s platelet count remained at 9000/μL on the 4th day after discontinuation of rifampicin. These results suggest that rifampicin was not the primary cause of drug-induced immune thrombocytopenia.

According to an approach proposed by Arnold et al[13], the diagnosis of drug-induced immune thrombocytopenia is based on the following four criteria: (1) Severity of thrombocytopenia: platelet count nadir below 20000/μL; (2) Clinical signs: Any bleeding; (3) Time to onset: Platelet counts fall 5-10 d after initiation of a new drug or exposure to a drug previously taken; and (4) Use of drugs already identified as responsible for drug-induced immune thrombocytopenia (with clinical and laboratory...
Figure 1 Changes of white blood cell, hemoglobin, platelet count, and medication during hospitalization. A: White blood cell count during hospitalization. B: Hemoglobin count during hospitalization; and C: Platelet count and timing of medications during hospitalization. LMOX: Latamoxef; RD: Rifampicin; INH: Isoniazid; EB: Ethambutol; PZA: Pyrazinamide; LEV: Levofoxacin; rhTPO: Recombinant human thrombopoietin; IVIG: Intravenous immunoglobulin, DXM: Dexamethasone, VK1: Vitamin K1.

Zhang RY et al. Severe thrombocytopenia induced by latamoxef

DOI: 10.12998/wjcc.v10.i22.7906 Copyright ©The Author(s) 2022.

tests), with the drug previously associating with drug-induced immune thrombocytopenia by clinical and laboratory criteria[13]. The first three criteria matched our patient’s presentation. Because of our hospital's limited laboratory conditions, we could not directly measure drug-dependent platelet antibodies using immunoassay or flow cytometry. Therefore, the fourth criterion could not be confirmed.

We excluded possible causes of thrombocytopenia such as tuberculosis of the spleen, pseudothrombocytopenia, primary immune thrombocytopenia, other drug-induced immune thrombocytopenia, food and beverages, infections, hypersplenism due to chronic liver disease, excessive alcohol intake, nutritional deficiencies, rheumatologic diseases, thrombotic microangiopathy, myelodyplasia, cancer with disseminated intravascular coagulation, cancer with bone marrow infiltration or suppression, and post-transfusion purpura. On the Naranjo scale, our patient scored six, placing him in the category of potential drug-related toxicity[14]. We could not rechallenge the patient with latamoxef for apparent reasons. According to our findings, latamoxef was the cause of the drug-induced immune thrombocytopenia.

The original instructions for latamoxef did not mention thrombocytopenia or coagulation dysfunction. Some studies mentioned that the N-methyl tetrazolium side-chain in latamoxef could lead to prothrombin deficiency, thrombocytopenia, platelet dysfunction, and bleeding. In such cases, one should supplement with vitamin K to reduce adverse reactions such as coagulation dysfunction and bleeding[2,15]. We searched PubMed, Embase, CNKI, Wan-Fang, and VIP database, and located four articles related to thrombocytopenia caused by latamoxef[5-7,16]. Although several studies reported that latamoxef could cause thrombocytopenia, thrombocytopenia induced by latamoxef in the Chinese population has never been reported previously. The literature suggests that one should use latamoxef cautiously in elderly patients with hepatic and renal dysfunction, history of ulcers, long-term use of broad-spectrum antibiotics, poor coagulation function, bleeding tendency, or use of anticoagulant and antiplatelet drugs[5,6,16]. The patient in our case had none of these risk factors; however, he had recurrent fevers for more than one month. Fever leads to high metabolic rates, and disseminated
tuberculosis is a consumptive disease that reduces immunity. He also had Crohn’s disease for more than five years and was treated with adalimumab as immunosuppressive therapy. Overall, the patient’s tolerance to drug-induced thrombocytopenia was lower than that of healthy adults. Therefore, we suggested that latamoxef should be discontinued immediately when patients with thrombocytopenia suspected to be caused by latamoxef, the platelet count is less than 20000/μL and complicated by bleeding or blood loss anemia. Moreover, first-line drug treatment such as corticosteroid, human immunoglobulin, platelet-raising drugs, and transfusion of platelets or coagulation factor should be considered to alleviate the symptoms as soon as possible. We also recommend that thrombocytopenia be included among the adverse effects in the Chinese instructions for latamoxef.

CONCLUSION

This is the first case of severe thrombocytopenia induced by latamoxef in a young Chinese patient. For patients treated with latamoxef, platelet counts should be carefully monitored, and clinicians should be vigilant for the appearance of scattered ecchymoses. Clinicians should discontinue latamoxef immediately when thrombocytopenia occurs in the context of latamoxef treatment, especially for patients with tuberculosis, malnutrition, polypharmacy, and immunosuppressive states, all of which are potential predisposing factors.

ACKNOWLEDGEMENTS

We would like to thank all medical staff who provided data and supported the study.

FOOTNOTES

Author contributions: Zhang RY proposed and supervised the study; Zhang JJ assisted with data analysis, Li JM, Xu YY and Xu YH managed the patient and collected samples; Cai XJ evaluated data and modified the manuscript; all authors contributed to the design and interpretation of the study and to further drafts.

Supported by the Special Research Fund of Hospital Pharmacy of Zhejiang Pharmaceutical Society, No. 2019ZYY27; and Zhejiang Medical and Health Science and Technology Plan, No. 2020KY741 and No. 2021KY910.

Informed consent statement: Informed written consent was obtained from the patient for publication of this report and any accompanying images.

Conflict-of-interest statement: All the authors report no relevant conflicts of interest for this article.

CARE Checklist (2016) statement: The authors have read the CARE Checklist (2016), and the manuscript was prepared and revised according to the CARE Checklist (2016).
Severe thrombocytopenia induced by latamoxef

Zhang RY et al.

REFERENCES

12. Sun P. Study on the mechanism of rifampicin-induced immune thrombocytopenia. Liaoning Province: Jinzhou Medical University, 2016