REVIEW

**E. coli** O104:H4 outbreak and haemolytic–uraemic syndrome

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Abstract

Background: The first cases of the European epidemic of Shiga toxin-producing *Escherichia coli* O104:H4 (STEC-O104:H4) infection were reported in Germany in April 2011.

Objectives: To characterize the 2011 STEC-O104:H4 outbreak and its management. A literature review is made to assess the state of the art in STEC–haemolytic–uraemic syndrome (HUS) epidemiology, pathogenesis, management and prognosis, focusing on critically ill adults.

Methods: References were obtained from the European Center for Disease Control and World Health Organization epidemiological updates, in addition to a PubMed search covering the period from 1980 to August 2011, including all published work on STEC-O104:H4 and reviews on HUS management and prognosis.

Results: The epidemic originated from a bean and seed sprouts farm in Lower Saxony, and was caused by the O104:H4 strain – a highly antibiotic resistant, hybrid enteroaggregative – Shiga toxin producing *E. coli* strain (STEC). The infection was characterized by increased HUS (25%) and a higher mortality rate. STEC enteritis and HUS are associated with significant mortality and morbidity, especially amongst patients with severe renal and neurological disorders. Management should center on prompt kidney protection by maintaining adequate renal perfusion, in addition to avoiding diuretics and nephrotoxic agents.

Conclusions: The published studies regarding antibiotic treatment lack good quality evidence. However, recent data suggest a potential modulating effect that explains the conflicting data but moreover suggest that azithromycin might be of use. Neutralizing monoclonal antibodies are a promising new therapy for STEC–HUS, with currently ongoing studies. Other treatments have not been shown to be superior to supportive therapy alone.

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Brote de E. coli O104:H4 y síndrome hemolítico-urémico

Resumen

Antecedentes: Los primeros casos de la epidemia europea de infección por E. coli productora de la toxina Shiga (STEC-O104:H4) se detectaron en Alemania en abril de 2011.

Objetivos: Caracterizar el brote de STEC-O104:H4 de 2011 y su gestión. Se realiza una revisión bibliográfica para evaluar el estado del arte en epidemiología, patogenia, tratamiento y pronóstico de la STEC-síndrome urémico-hemolítico (SHU), con especial hincapié en los adultos críticamente enfermos.

Métodos: Se obtuvieron referencias de las actualizaciones epidemiológicas del Centro Europeo de Control de Enfermedades y de la Organización Mundial de la Salud, además de una búsqueda en PubMed relativa al periodo de 1980 a agosto de 2011, incluidas todas las publicaciones sobre la STEC-O104:H4 y las revisiones sobre el tratamiento y el pronóstico del SHU.

Resultados: La epidemia tuvo su origen en unos brotes germinados de una plantación de Baja Sajonia y estuvo causada por la cepa O104:H4, una cepa híbrida de E. coli enteragregatriz productora de la toxina Shiga (STEC) con una elevada resistencia a los antibióticos. La infección se caracterizó por un aumento del SHU (25%) y por una tasa de mortalidad más elevada. El SHU y la enteritis de la STEC se asocian a una morbimortalidad importante, especialmente en pacientes con trastornos neurológicos y renales graves. El tratamiento debe centrarse en una rápida protección renal, manteniendo una perfusión renal adecuada, además de evitar los diuréticos y los agentes nefrotóxicos.

Conclusiones: Los estudios publicados en relación con el tratamiento antibiótico no están respaldados por datos de calidad. Sin embargo, los datos recientes sugieren un potencial efecto modulador que no solo explica la incoherencia en los datos sino que además sugiere que la azitromicina podría resultar de utilidad. Los anticuerpos monoclonales neutralizadores son un prometedor tratamiento nuevo para la STEC-SHU, y actualmente se están llevando a cabo estudios sobre ellos. Otros tratamientos no han resultado ser superiores a la administración de cuidados paliativos exclusivamente.

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E. coli, a major facultative inhabitant of the large intestine, is a Gram-negative bacilli and facultatively anaerobic. Strains are classified according to the O (lipopolysaccharide) and H (flagellum) antigens. Enterohaemorrhagic (EHEC) strains are characterized by its genomic and plasmidic virulence factors. The first one translates into the production of one or two types of Shiga-like toxins (STX), STX-1 and STX-2. Whereas the latter endows it with genes encoding proteins key to its virulence and pathogenesis: such as catalase–peroxidase enzymes, haemolysins, proteases, secretion proteins and anticoagulants. Characterized by a low infectious dose (10–100 microorganisms), it causes haemorrhagic colitis that can be complicated with HUS. Strains associated with HUS usually harbour STX-2, curiously, it is less frequent amongst strains with both STX genes. The most common serotype of outbreaks and HUS is the STEC-O157:H7. Whilst enteroaggregative E. coli presents with persistent watery diarrhoea, usually in children. Infection with EHEC is commonly associated with contaminated beef; however, a retrospective review of 350 outbreaks revealed that even though gastrointestinal tract of cattle is the most important reservoir and food faecally contaminated by cattle (including raw milk) is the first mechanism of transmission (52% of cases), beef is a less frequent vehicle. Foodborne transmission is followed frequently by person-to-person (14%) and waterborne, including swimming (9%). Person-to-person transmission peaks during diarrhoea; post-symptomatic shedding can occur.

**Pathogenesis and clinical presentation**

After its adherence to enterocytes, STEC produces attaching-effacing lesions, through its plasmidic encoded proteins. Providing an entry point into the intravascular space for toxins. Targeting endothelial cells, STX halt cell protein synthesis causing cell death. Which translates into vascular damage; predominantly locally (at the intestine) and at the kidneys. Producing variable prothrombotic states in addition to non-inflammatory bloody diarrhoea and kidney failure. Symptoms usually present after 72 h of ingestion of the inoculum. Initially, watery diarrhoea is accompanied by abdominal pain, vomiting and usually without fever (although almost 50% of patients report fever during the prodromic phase), 48–72 h later, haemorrhagic enterocolitis ensues in the 90% of cases. Improvement of diarrhoea occurs at the 7th day of symptoms with spontaneous resolution in the 85% of cases, whilst 15% develop HUS. Post-symptomatic shedding can be present from the 3rd to the 7th day of initiation of symptoms (Fig. 2).

Overall, HUS’s incidence in STEC infections rounds 6–9% and in children 15%; whilst, in the STEC-O104:H4 outbreak it rose to 25%. Conversely, 70% of HUS are caused by EHEC; other causes include pneumococcal pneumonia and complement inherited-abnormalities. HUS is defined as microangiopathic haemolytic anaemia, thrombocytopenia and acute kidney injury (AKI). Despite its low mortality (5–7%), it is associated to significant morbidity: 50% of patients require extra-renal depurative techniques and 5–10% have renal sequelae. Initially considered as a part of the spectrum of the same thrombotic non-vasculitic disorder, manifesting either with thrombotic thrombocytopenic purpura (TTP) in adults and HUS in children. Actually a distinctive pathophysiology is known in each case; TTP is an autoimmune disorder caused by inhibition of Von Willebrand factor, whilst HUS is characterized by fibrinolysis inhibition, increased fibrin production and thrombin formation, entirely secondary to STX injury and almost never manifesting with bacteraemia. The core pathogenesis of HUS is the formation of microvascular thrombi, followed by endothelial oedema and vascular leakage. New findings suggest that STX also directly interferes with the alternative complement pathway and that it is a key pathogenic mechanism. Thrombocytopenia is the first abnormality in
E. coli O104:H4 outbreak and haemolytic–uraemic syndrome

Figure 2 Pathogenesis of E. coli enteritis and haemolytic–uraemic syndrome. AKI: acute kidney injury; CNS: central nervous system. HUS triad: thrombocytopenia, anaemia and AKI.

STEC infections, either the patient develops HUS or not, and lasts up to one week. It is secondary to thromboctye consumption and somewhat to voluminous expansion. A reliable parameter of clinical resolution, platelet count with stable or ascending trends translates into improvement and in most cases presents the possibility of discharge. If renal failure has not developed up to this point, almost it unequivocally will not occur afterwards. Microangiopathic haemolytic anaemia is a result of the traumatic rupture due to endothelial thrombi and is the most persistent abnormality, usually preceding renal failure. Azotemia and acute tubular necrosis are responsible for AKI. Overload is due to oliguria and fluid resuscitation. Neurological complications of HUS are important determinant of morbidity and the most common cause of mortality. It encompasses a wide array of clinical findings, from minor changes in mental status to major neurologic deficits such as seizures, coma and hemiparesis. Proposed mechanisms include thrombotic occlusions (coagulative necrosis due to microthrombosis without haemorrhage), microangiopathic changes of small vessels (oedema and focal haemorrhage), direct injury of toxin, ischaemia–reperfusion injury, hypertension and metabolic derangement (hypoglycaemia, dehydration, acid–base disturbances, changes in serum osmolality or azotemia). Neurological damage occurs in 50–100% of children with HUS, the most frequent presentation is a generalized or partial seizures, followed by coma, irritability and focal signs (hemiparesis or aphasia). Neurological sequelae follow in most cases. Diffuse encephalopathy and SWCP score are useful predictors of bad neurological outcome. Neuroimaging findings include ischaemia and are usually located at basal ganglia and thalamus.

Both watery and bloody diarrhoea are probably due to intestinal ischaemia, rather than by invasive/inflammatory damage, and can evolve to necrosis and perforation. Elevations in serum amylase and lipase have been reported, without clinical implications. Heart failure is usually secondary to volume overload; however, elevation of cardiac markers such as troponins suggests myocardial infarction rather than reduced renal clearance.

Workup

Sorbitol-MacConkey agar cultures allow STEC-O157:H7 serotype identification. The highest likelihood of positive culture occurs on days 1–6 from onset of diarrhoea; usually positive on the third day. Molecular diagnosis of STEC can be done in stool directly with the detection of STX with enzyme linked immunosorbent assay (ELISA) or indirectly (STX genes) with polymerase chain reaction (PCR). Specific serogroup identification can be achieved using ELISA and Western blot with specific antibodies detection, both in the acute and convalescent periods.

Management

Most efforts are directed towards kidney protection. Maintaining an appropriate renal perfusion is key in preventing HUS. Studies show that adequate pre-HUS volume expansion is associated with lesser AKI in HUS. Therefore, therapies that could deplete the intravascular volume, such as diuretics and medication that could elevate the risk or worsen AKI (NSAIDs, ACE inhibitors and nephrotoxic agents) should be avoided. Renal replacement therapy should be implemented according to the usual indications in ARF. Careful fluid balance should be sought, taking into account the increased risk for third-space liquid accumulation secondary to vascular leakage and hypoalbuminaemia. Hypertension is preferred to be treated with vasodilators. Erythrocyte transfusions also should follow standard indications. Anti-motility agents and narcotics are not recommended, although is not clear whether they increase the risk of HUS and neurological complications or not.

Ineffective interventions include those directed to equilibrate the prothrombotic state and to the inhibition of STX and auto-immune mechanisms. Anticoagulation, anti-aggregation and fibrinolysis are not associated with better outcome, instead, they highly increase risk of bleeding (mainly intracranial bleeding); compared to supportive therapy, OR for bleeding is 25.89 (95% CI 3.67–182.83). Corticosteroids, plasmapheresis and plasma infusion are not
superior to supportive therapy, which is logical considering the absence of immune-mediated pathways. Despite that SHU is mainly a consequence of direct action of the Shiga-toxin, binding agents also fail to contribute to its management, which can be explained because at the time of SHU onset, STX-endothelial damage has been already established.\(^1\)\(^8\)

Currently much controversy revolves around antibiotic treatment of EHEC. Data from a retrospective study of 1993’s USA \(E.\ coli\) O157:H7 outbreak reported that antibiotic administration was not associated with a decreased risk for HUS development,\(^6\) setting a trend of not treating with antibiotics for these infections. Proposed mechanisms included that bacterial lysis caused liberation of STX and the induction of bacteriophages (where STX genes are located), which further increased STX levels. It also caused selection of resistant strains. Considering that the main pathogenic factor is direct toxin injury and that at the time of onset of diarrhoea, the STX is already in circulation; it suggested that little benefit would come from administration of antibiotics.\(^7\)

Thereafter, on 2003, a meta-analysis\(^9\) that included 9 clinical studies, of which 6 retrospective, suggested that antibiotic treatment with fosfomycin might be a protective factor for HUS in \(E.\ coli\) O157:H7 enteritis, even though studies lacked adequate methodology to support the evidence. The pooled OR did not show a clear result to favour any trend and conclusions were based on a pooled-OR that excluded two studies that clearly associated antibiotic treatment with increased risk of HUS. Amongst the pro-antibiotic treatment studies, the Ikeda et al.,\(^10\) a prospective cohort, evaluated 36 cases of patients receiving fosfomycin vs. other antibiotic (not specified), OR was a result of a univariate analysis that yielded 0.12 (95% CI 0.02–0.74). Cimolai et al.\(^11\) published a retrospective case-control, including 27 cases reporting an OR: 0.63 (95% CI 0.27–1.48); however, neither the type of antibiotic nor the diarrhoea-antibiotic administration interval was stated. On the other hand, one of the excluded studies, Wong et al.,\(^12\) a prospective cohort that included 10 cases in children (<10 years) where antibiotics (trimethoprim-sulfamethoxazole, amoxicillin and cephalosporins) were administered within 3 days of diarrhoea onset (DO), yielded an adjusted OR for HUS of 17.9 with statistical significance (unadjusted OR: 14.25; 95% CI 3.62–56.1). The other excluded study was the Pavia et al.,\(^13\) a retrospective case-control and randomized trial of 8 cases which received antibiotics within 3 days of DO; with a OR for HUS of 80.6 with statistical significance (95% CI 3.38–1922).

Lynn et al.\(^14\) published a 4 years prospective surveillance of childhood HUS in United Kingdom and Ireland; including 413 cases of HUS. Statistical significance was found between neither mortality and antibiotic treatment, nor antibiotic treatment with ciprofloxacin vs. metronidazole/penicillin/2nd–3rd generation cephalosporin.

On 2006 a systematic review\(^15\) that included all cases published between years 1982 and 2005, analysed whether antibiotics are beneficial or detrimental for \(E.\ coli\) O157:H7 infection. In total, 63 in vitro and clinical studies were included, case-reports were excluded. Of the experimental data, they found that: (a) bacteria can recover and continue producing extracellular toxins after exposure to continuously changing sub-lethal antibiotics concentrations; (b) antibiotics might be detrimental, given that increased release of toxins was found in cultures exposed to sub-inhibitory and sub-lethal concentrations of polymyxin B, trimethoprim-sulfamethoxazole, ciprofloxacin, cefixime and tetracycline; and (c) fewer STX is released when exposed to sub-minimum inhibitory concentrations of roxithromycin, rokitamycin and clindamycin. Concluding that sub-lethal concentrations of antibiotics might have a modulating effect on STX production. Within the clinical studies that advocate to the beneficial effect of antibiotic treatment that were not included in the 2003 meta-analysis were the Martin et al.,\(^16\) a retrospective cohort of 117 patients (<18 years), of which only 20 were treated with antibiotics during prodromal illness (without specification) showed a mild clinical course and good outcome; it also reported that antibiotic treatment was less common in the group of severe outcome. Higami et al.\(^17\) conducted a retrospective control including 216 children, comparing antibiotic regimens at the 3rd day of OD, less HUS incidence was found in the group treated with new quinolones in comparison to cephalosporins. Shiomi et al.,\(^18\) with a retrospective control study of 42 children, found that antibiotic treatment on 3rd day of OD with oral fluoroquinolines (FQ) prevents HUS when compared to a endovenous FQ plus cefotaxime regime. On the other hand, studies exposing a detrimental effect of antibiotics included the Slutsker et al.,\(^19\) a retrospective case-control comprising 230 patients of all ages, showed that sulfamethoxazole given at the 3rd day of OD was related with an OR for HUS 11.5%, in the subgroup of <13 years. Carter et al.,\(^20\) through a retrospective case-control of 73 patients also of all ages; analysed timing of initiation of antibiotics; when given before the onset of symptoms (OOS) increased secondary person to person infection was reported; whereas when given after OOS, elevated case-fatality rate was found, probably due to disease severity bias. Ostroff et al.,\(^21\) in a retrospective case-control of 80 patients (including all ages), compared TMP/SMX with gentamicin initiated at the 4th day; TMP/SMX showed a OR for HUS: 3.1 (even though the CI included 1); and gentamicin, OR: 9.1 (CI = 1). Finally, Dundas et al.,\(^22\) in a retrospective cohort of 120 patients of all ages, analysed those who received antibiotic 4 weeks before OOS and found association with increased incidence of HUS, except for ciprofloxacin that did not show any difference compared to controls; no association with mortality was found.

Amongst the latest evidence addressing this issue, despite its pure experimental nature, provides a coherent explanation that encompasses the evidence found out about antibiotics and STX modulation until now.\(^23\) It studied different \(E.\ coli\) O157:H7 strains, different plasmids and different antibiotics. Results showed that phages can infect harmless intestinal \(E.\ coli\), recruiting them to produce STX. Grow-inhibitory levels of antibiotics decrease STX production. Even when O157:H7’s viability remained high, azithromycin significantly reduced STX production. Nonetheless, sub-Inhibitory levels that target DNA synthesis (ciprofloxacin/TMP-SMX) increase STX production; whilst ciprofloxacin’s complete suppression of O157:H7 markedly increases levels of STX. Antibiotics targeting cell wall,
transcription and translation do not have any effect on STX levels. Based on the evidence, during this outbreak Germany’s infectious diseases society recommended to avoid the use of FQ, TMP-SMX, aminoglycosides and fosfomycin in EHEC patients. Table 1 summarizes these studies according to the relationship between HUS and antibiotic treatment.

Neutralizing monoclonal antibody of STX is promising amongst new therapies in the development of Shiga toxin. Eculizumab is a recombinant, humanized, monoclonal immunoglobulin G antibody derived from murine myeloma cells, directed against protein C5 inhibiting activation of the terminal complement pathway. Authorized only for paroxysmal nocturnal haemoglobinuria; in this setting, reduces intravascular haemolysis, anaemia and thrombotic complications. However, since terminal complement is necessary to prevent and limit a thrombotic complications. However, results are limited by the design of the studies. However, results are limited by the design of the studies.

### Table 1

<table>
<thead>
<tr>
<th>Study (first author)</th>
<th>Type of study</th>
<th>Sample size (number of patients)</th>
<th>OR (95% CI)</th>
</tr>
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<tr>
<td>Safdar et al.¹⁹</td>
<td>Meta-analysis</td>
<td>a</td>
<td>0.83 (0.54–1.26)³⁶</td>
</tr>
<tr>
<td>Ikeda et al.²⁰</td>
<td>Prospective cohort</td>
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<td>0.12 (0.02–0.75)</td>
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<tr>
<td>Cimolai et al.²¹</td>
<td>Retrospective case-control</td>
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<td>0.63 (0.27–1.43)</td>
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<tr>
<td>Dundas et al.²²</td>
<td>Retrospective cohort</td>
<td>120</td>
<td>1.23 (0.43–3.8)</td>
</tr>
<tr>
<td>Martin et al.²⁶</td>
<td>Retrospective cohort</td>
<td>117</td>
<td>9.4 (1.2–199)</td>
</tr>
<tr>
<td>Higami et al.²⁷</td>
<td>Retrospective control</td>
<td>216</td>
<td>NS³</td>
</tr>
<tr>
<td>Shiomi et al.²⁸</td>
<td>Retrospective control</td>
<td>42</td>
<td>NS³</td>
</tr>
</tbody>
</table>

### Prognosis

Up to 10% of patients who suffered HUS, wind up with renal sequels. In a meta-analysis/meta-regression of 49 studies from 1950 to 2001 that included 3,476 patients aging from 1 month to 18 years from 18 countries, with a follow-up of 4.4 years, confirmed that death or end-stage renal disease (ERSD) occurs in approximately 12% with EHEC-HUS and that one quarter of survivors suffer long-term renal disease. Worse prognosis is associated with severity of acute illness, especially with early requirement of renal depurative techniques and CNS severe manifestations.

### Conclusions

E. coli O104:H4 strain is characterized by a low infection dose, hybrid adhesion proteins and antibiotic resistance (ESBL). 2011’s outbreak presented with increased HUS and greater fatality rate. Management should focus on prompt kidney protection by maintaining adequate renal perfusion in addition to avoidance of diuretics and nephrotoxic agents. Antibiotics have potential modulator effect, of which azithromycin might be of use for STEC-HUS treatment. However, results are limited by the design of the
selected studies: case–control, not randomized-controlled with high heterogeneity and lacking information of both sample and treatment. Well-designed studies are needed to establish the role of antibiotics in this setting. Neutralizing monoclonal antibodies are a promising new therapy for STEC–HUS, with currently ongoing studies. Platelet trend is a reliable parameter for management guidance. Other therapies have not proven to be superior to supportive therapy alone.

Conflicts of interest

Authors do not have any financial relationship with a commercial entity that has an interest in the subject of this manuscript.

References


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