

**CLINICAL MICROBIOLOGY AND INFECTION**

VOLUME 20, SUPPLEMENT 2, MARCH 2014

**European Society of Clinical Microbiology and  
Infectious Diseases: Update of the Treatment  
Guidance Document for *Clostridium difficile*  
Infection**

Publication of this supplement was commissioned and funded by ESCMID



**WILEY**  
Blackwell

# Clinical Microbiology and Infection

VOLUME 20, SUPPLEMENT 2, MARCH 2014

- Original Article*      1    European Society of Clinical Microbiology and Infectious Diseases:  
Update of the Treatment Guidance Document for *Clostridium difficile*  
Infection  
*S. B. Debast, M. P. Bauer and E. J. Kuijper on behalf of the Committee*

# European Society of Clinical Microbiology and Infectious Diseases: update of the treatment guidance document for *Clostridium difficile* infection

S. B. Debast<sup>1</sup>, M. P. Bauer<sup>2</sup>, E. J. Kuijper<sup>3</sup>, on behalf of the Committee\*

1) Department of Medical Microbiology, Radboud University Medical Center, Nijmegen, Departments of 2) Infectious Diseases and 3) Medical Microbiology, Centre for Infectious Diseases, Leiden University Medical Centre, Leiden, the Netherlands

## Abstract

In 2009 the first European Society of Clinical Microbiology and Infection (ESCMID) treatment guidance document for *Clostridium difficile* infection (CDI) was published. The guideline has been applied widely in clinical practice. In this document an update and review on the comparative effectiveness of the currently available treatment modalities of CDI is given, thereby providing evidence-based recommendations on this issue. A computerized literature search was carried out to investigate randomized and non-randomized trials investigating the effect of an intervention on the clinical outcome of CDI. The Grades of Recommendation Assessment, Development and Evaluation (GRADE) system was used to grade the strength of our recommendations and the quality of the evidence. The ESCMID and an international team of experts from 11 European countries supported the process. To improve clinical guidance in the treatment of CDI, recommendations are specified for various patient groups, e.g. initial non-severe disease, severe CDI, first recurrence or risk for recurrent disease, multiple recurrences and treatment of CDI when oral administration is not possible. Treatment options that are reviewed include: antibiotics, toxin-binding resins and polymers, immunotherapy, probiotics, and faecal or bacterial intestinal transplantation. Except for very mild CDI that is clearly induced by antibiotic usage antibiotic treatment is advised. The main antibiotics that are recommended are metronidazole, vancomycin and fidaxomicin. Faecal transplantation is strongly recommended for multiple recurrent CDI. In case of perforation of the colon and/or systemic inflammation and deteriorating clinical condition despite antibiotic therapy, total abdominal colectomy or diverting loop ileostomy combined with colonic lavage is recommended.

**Keywords:** *Clostridium difficile* infection, guideline, recommendations, review, treatment

**Original Submission:** 23 July 2013; **Revised Submission:** 22 September 2013; **Accepted:** 27 September 2013

Editor: M. Paul

**Article published online:** 5 October 2013

*Clin Microbiol Infect* 2014; **20** (Suppl. 2): 1–26

**Corresponding author:** E. J. Kuijper, Department of Medical Microbiology, Centre for Infectious Disease, Leiden University Medical Centre, PO Box 9600, 2300 RC Leiden, the Netherlands  
**E-mail:** e.j.kuijper@lumc.nl

\*Committee details see Appendix 1.

## Introduction

The previous European Society of Clinical Microbiology and Infection (ESCMID) guidance document, which has been applied widely in clinical practice, dates from 2009 [1]. Meanwhile, new treatments for *Clostridium difficile* infection (CDI) have been developed and limitations of the currently recommended

treatment options of CDI are considered. As the current ESCMID treatment guidance document is already implemented in clinical practice, an update of this widely applied guidance document is essential to further improve uniformity of national hospital infection treatment policies for CDI in Europe. In particular, after the recent development of new alternative drugs for the treatment of CDI (e.g. fidaxomicin) in the USA and Europe, there has been an increasing need for an update on the comparative effectiveness of the currently available antibiotic agents in the treatment of CDI, thereby providing evidence-based recommendations on this issue.

The objectives of this document are to:

1. Provide an overview of currently available CDI treatment options
2. Develop an evidence-based update of treatment recommendations

## Update Methodology

Studies on CDI treatment were found with a computerized literature search of PUBMED and Google Scholar using the terms 'Clostridium difficile AND (treatment OR trial)'. All randomized and non-randomized trials investigating the effect of an intervention on the clinical outcome (resolution or recurrence of diarrhoea; incidence of complications) of CDI published in any language were included. Studies investigating carriage or other purely microbiological parameters were not considered sufficient evidence for treatment strategies. The resulting literature from 1978 was reviewed and analysed. Furthermore, systematic reviews from the most recent Cochrane analysis [2] and the up-dated guidelines of the Infectious Diseases Society of America, the Australasian Society for Infectious Diseases, the American College of Gastroenterology, and the Health Protection Agency/Public Health England guidance document (<http://www.hpa.org.uk>) were evaluated [3–5]. Recommendations were based on a systematic assessment of the quality of evidence. The Grades of Recommendation Assessment, Development and Evaluation (GRADE) system was used to grade the strength of our recommendations and the quality of the evidence [6,7].

Draft versions of the guideline were written by the executive committee (consisting of: S. Debast, M. Bauer and E. Kuijper) and criticized by the Executive Committee and advisors. After this, consensus was reached, resulting in the final version. The methods to evaluate the quality of evidence and to reach group consensus recommendations were based on the method described by Ullmann *et al.* [8].

Definition of the strength of recommendation is given in Table 1. The quality of the published evidence is defined in Table 2a. Grouping quality of evidence into three levels only may lead to diverse types of published evidence being assigned specifically to a level II. To increase transparency in the evaluation of the evidence an index (Table 2b) to the level II recommendations was added where appropriate.

The guideline followed the Appraisal of Guidelines Research and Evaluation Collaboration (AGREE) self-assessment tool [9].

**TABLE 1.** Definition of the Strength of Recommendation Grade (SoR) ESCMID (adapted from ref. [8])

Strength	Definition
A	Strongly supports a recommendation for use
B	Moderately supports a recommendation for use
C	Marginally supports a recommendation for use
D	Supports a recommendation AGAINST use

## Definitions

### Diagnosis

The diagnosis of CDI is based on (I) a combination of signs and symptoms, confirmed by microbiological evidence of *C. difficile* toxin and toxin-producing *C. difficile* in stools, in the absence of another cause, or (ii) colonoscopic or histopathological findings demonstrating pseudomembranous colitis [1,3,10–12].

There are many different approaches that can be used in the laboratory diagnosis of CDI; however, the best standard laboratory test for diagnosis has not been established. Diagnostic tests for CDI include: (i) detection of *C. difficile* products: cell culture cytotoxicity assay (CCA), glutamate dehydrogenase (GDH) and Toxins A and/or B, (ii) toxigenic culture of *C. difficile*, and (iii) nucleic acid amplification tests (NAAT): 16S RNA, toxin genes, GDH genes. Preferably a two- or three-stage algorithm is performed to diagnose CDI, in which a positive first test is confirmed with one or two confirmatory tests or a reference method [3,4,12,13]. Faeces samples could be investigated with an enzyme immunoassay detecting GDH, an enzyme immunoassay detecting toxins A and B, or NAAT detecting Toxin B (TcdB). Samples with a negative test result can be reported as negative. Faeces samples with a positive first test result should be re-tested with a method to detect free faeces toxins, or with a method to detect GDH or toxin genes, dependent on the assay applied as first screening test. If free faeces toxins are absent but *C. difficile* TcdB gene or GDH are present, CDI cannot be differentiated from asymptomatic colonization. Recently, a large study was presented in which several diagnostic algorithms were evaluated to optimize the laboratory diagnosis of CDI [14]. The investigators concluded that two-stage algorithms improve diagnosis of CDI. Two commonly recommended methods in the laboratory diagnosis of CDI are the use of GDH detection in stools as a means of screening for CDI, confirmed by NAAT such as PCR to detect toxigenic strains of *C. difficile* [4,12]. Furthermore, patients with a positive stool toxin had *C. difficile* disease with an increased risk of mortality compared with patients with only a positive toxigenic culture, thereby implying that stool toxin testing should be included in a testing algorithm to optimize *C. difficile* diagnostic testing [15].

Diarrhoea is defined as loose stools, i.e. taking the shape of the receptacle or corresponding to Bristol stool chart types 5–7, plus a stool frequency of three stools in 24 or fewer consecutive hours or more frequently than is normal for the individual (definition World Health Organization, <http://www.who.int/topics/diarrhoea>) [1,3,16–18].

Clinical pictures compatible with CDI are summarized in Table 3.

**TABLE 2.** Definition of the Quality of Evidence (QoE) ESCMID. Adapted from ref. [8]

Quality of evidence	Definition
2a: Level I	Evidence from at least one properly designed randomized, controlled trial.
II	Evidence from at least one well-designed clinical trial, without randomization; from cohort or case-control analytic studies (preferably from more than one centre); from multiple time series; or from dramatic results of uncontrolled experiments.
III	Evidence from opinions of respected authorities, based on clinical experience, descriptive case studies, or reports of expert committees.
2b: Index	
r	Meta-analysis or systematic review of randomized controlled trials.
t	Transferred evidence, i.e. results from different patient cohorts, or similar immune-status situation.
h	Comparator group is a historical control.
u	Uncontrolled trial.
a	Abstract or poster of a study published at an international meeting.

**Definition of *Clostridium difficile* infection.** An episode of CDI is defined as:

A clinical picture compatible with CDI and microbiological evidence of free toxins and the presence of *C. difficile* in stool without reasonable evidence of another cause of diarrhoea.

**or**

Pseudomembranous colitis as diagnosed during endoscopy, after colectomy or on autopsy [3,11,19].

### Treatment response

**Definition of treatment response.** Treatment response is present when either stool frequency decreases or stool consistency improves and parameters of disease severity (clinical, laboratory, radiological) improve and no new signs of severe disease develop. In all other cases, treatment is considered a failure. Treatment response should be observed daily and evaluated after at least 3 days, assuming that the patient is not worsening on treatment. Treatment with metronidazole, in particular, may result in a clinical response only after 3–5 days [21–23]. After clinical response, it may take weeks for stool consistency and frequency to become entirely normal [23,24].

### Recurrences

**Definition of recurrent *Clostridium difficile* infection.** Recurrence is present when CDI re-occurs within 8 weeks after the onset of a previous episode, provided the symptoms from the previous episode resolved after completion of initial treatment [4,11].

It is not feasible to distinguish recurrence due to relapse (renewed symptoms from already present CDI) from recurrence due to reinfection in daily practice [20,25–28].

**TABLE 3.** Clinical pictures compatible with *Clostridium difficile* infection. Adapted from refs [1,3,11,19,20]

Sign/symptom	Definition
Diarrhoea	Loose stools, i.e. taking the shape of the receptacle or corresponding to Bristol stool chart types 5–7, plus a stool frequency of three stools in 24 or fewer consecutive hours or more frequently than is normal for the individual.
Ileus	Signs of severely disturbed bowel function such as vomiting and absence of stool with radiological signs of bowel distension.
Toxic megacolon	Radiological signs of distension of the colon (>6 cm in transverse width of colon) and signs of a severe systemic inflammatory response.

### Severity of disease

**Definition of severe *Clostridium difficile* infection.** Severe CDI is defined as an episode of CDI with (one or more specific signs and symptoms of) severe colitis or a complicated course of disease, with significant systemic toxin effects and shock, resulting in need for ICU admission, colectomy or death [1,4,29].

*Clostridium difficile* infection without signs of severe colitis in patients with greater age ( $\geq 65$  years), serious comorbidity, Intensive Care Unit (ICU) admission, or immunodeficiency may also be considered at increased risk of severe CDI [30,31].

An overview of characteristics in patients with CDI that are assumed to correlate with the severity of colitis is given in Table 4 [32–39]. We must stress that the prognostic value of these markers is uncertain.

### Clinical prediction markers

**Evidence.** Clinical studies indicate superiority of specific treatment strategies depending on the severity of disease. In addition, alternative treatment options have been developed, that may be more effective in preventing recurrence of disease. Unfortunately some of the novel treatment strategies can be very expensive, and may only be cost-effective for a certain group of patients depending on the stage and severity of disease. This emphasizes the importance for better identification of clinical markers, preferably early in the course of disease, which might predict the benefit from specific treatment regimens to decrease CDI-related complications, mortality or recurrences. Surprisingly little prospective and validated research has been performed on clinical predictors of outcome [40]. Furthermore, for some complications of CDI, such as ICU admission or death, it is difficult to determine to what extent the complication can be attributed to CDI as opposed to the presenting acute illness(es) or comorbidities.

A wide variety of risk factors for severe or recurrent CDI have been suggested in literature, which makes it difficult to set a rigid clinical prediction rule [1,25,41–46]. Recently, a

**TABLE 4.** Patient characteristics that could reasonably be assumed to correlate positively with severity of colitis in the absence of another explanation for these findings

Category	Signs/symptoms
Physical examination	Fever (core body temperature >38.5°C). Rigors (uncontrollable shaking and a feeling of cold followed by a rise in body temperature). Haemodynamic instability including signs of distributive shock. Respiratory failure requiring mechanical ventilation. Signs and symptoms of peritonitis. Signs and symptoms of colonic ileus. Admixture of blood with stools is rare in <i>Clostridium difficile</i> infection (CDI) and the correlation with severity of disease is uncertain.
Laboratory investigations	Marked leucocytosis (leucocyte count >15 × 10 <sup>9</sup> /L). Marked left shift (band neutrophils >20% of leucocytes). Rise in serum creatinine (>50% above the baseline). Elevated serum lactate (≥5 mM). Markedly reduced serum albumin (<30 g/L). Pseudomembranous colitis.
Colonoscopy or sigmoidoscopy	There is insufficient knowledge on the correlation of endoscopic findings compatible with CDI, such as oedema, erythema, friability and ulceration, and the severity of disease.
Imaging	Distension of large intestine (>6 cm in transverse width of colon). Colonic wall thickening including low-attenuation mural thickening. Pericolonic fat stranding. Ascites not explained by other causes. The correlation of haustral or mucosal thickening, including thumbprinting, pseudopolyps and plaques, with severity of disease is unclear.

systematic review was performed to derive and validate clinical rules to predict recurrences, complications and mortality [46]. Most studies were found to have a high risk of bias because of small sample sizes and much heterogeneity in the variables used, except for leucocytosis, serum albumin and age [46]. Bauer *et al.* used a database of two randomized controlled trials, which contained information for a large patient group (1105 patients) with CDI, to investigate the prognostic value of three markers for severe CDI. They found that both leucocytosis and renal failure are useful predictors of a complicated course of CDI, if measured on the day of diagnosis [45].

A recent meta-analysis of two pivotal randomized controlled trials comparing fidaxomicin and vancomycin revealed previous vancomycin or metronidazole treatment in the 24 h before randomization, low eosinophil count (<0.1 × 10<sup>9</sup>/L) and low albumin level to be independent predictors of persistent diarrhoea or death in the first 12 days [40]. Recently Miller *et al.* [36] analysed the same two clinical therapeutic trials to derive and validate a categorization system to discriminate among CDI patients and correlate the grouping with treatment response. They concluded that a combination of five clinical and laboratory variables measured at the time of CDI diagnosis, combined into a scoring system, were able to

accurately predict treatment response to CDI therapy with fidaxomicin and vancomycin. These variables include: age, treatment with systemic antibiotics, leucocyte count, albumin and temperature (ATLAS).

Strain type has been suggested as an additional cause of excess morbidity, disease severity and higher recurrence rates of CDI. In a Canadian study [47], PCR ribotype 027 was correlated with more severe disease and fatal outcome among patients at almost all ages. Some studies on the other hand suggested that PCR ribotype 027 strains might only be associated with worse outcome in settings where 027 strains are epidemic, and not in an endemic situation [38,48]. However, these findings are questioned by others [49]. Recently, a large study by Walker *et al.* clearly showed that strain types varied in the overall impact on mortality and biomarkers (predominantly those associated with inflammatory pathways) [50]. Besides *C. difficile* PCR ribotype 027, other strains are also associated with outbreaks and severe *C. difficile* infection, e.g. PCR ribotype 078 [51]. Despite increased virulence of specific strain types, the value of the PCR ribotype as a prediction marker for disease severity may be limited, as the ribotype involved in an infection is commonly not known upon diagnosis. However, in an epidemic situation the PCR ribotype may be taken into account in deciding on the choice of empirical treatment regimens [21,39].

The level of host immune response to *C. difficile* exposure has been shown to be an important determinant of the severity and duration of clinical manifestations [52–57]. Anti-toxin antibody levels have been demonstrated to be higher in healthy adult controls compared with healthy children, and levels were found to fall with increasing age. In addition, anti-toxin antibodies increased after resolution of diarrhoea, which coincided with decreased incidence of CDI recurrence [57]. Inability to mount an adequate humoral immune response (e.g. during use of rituximab) may therefore be an important additional prediction marker for severe and/or recurrent CDI [25,57–62]. Unfortunately, in most cases this information is not available at presentation/diagnosis; also, as the strength of evidence for immunodeficiency as an independent predictor for severe and/or recurrent CDI is still limited, we did not include this risk factor as a separate prediction marker.

The results from individual studies, reviews and meta-analyses on prognostic markers for CDI were evaluated to reach a group consensus on a selection of markers that may be useful in clinical practice to distinguish patients with increased risk for severe or life-threatening CDI and recurrences. For detailed recommendations we refer to Tables 5 and 6.

**Recommendations.** *Clostridium difficile* infection is judged to be severe when one or more of the clinical markers of severe

**TABLE 5. Prognostic markers that can be used to determine (increased risk of developing) severe *Clostridium difficile* infection (CDI)**

Characteristics	SoR <sup>a</sup>	QoE	Ref (s)	Comment(s)
Age ( $\geq 65$ years)	A	IIr	[32,41,46]	Large cohort study on CDI mortality at 30 days, and review of studies of factors associated with CDI outcome [41]. Systematic review of studies describing the derivation or validation of Clinical Prediction Rules for unfavourable outcomes of CDI [46]: in general methodological biases and weak validities.
Marked leucocytosis (leucocyte count $> 15 \times 10^9/L$ )	A	IIrht	[32,37,39,45,46,63,64]	Systematic review [46]: in general methodological biases and weak validities. Cohort study: severity score on malignancy, white blood cell count, blood albumin, and creatinine [37]. Retrospective cohort study on risk factors for severe CDI: death $< 30$ days, ICU, colectomy or intestinal perforation [32].
Decreased blood albumin ( $< 30$ g/L)	A	IIr	[32,37,40,46,65]	Systematic review [46]: in general methodological biases and weak validities.
Rise in serum creatinine level ( $\geq 133$ $\mu M$ or $\geq 1.5$ times the premorbid level)	A	IIht	[32,37,41,45]	Depending on the timing of measurement around CDI diagnosis [45].
Comorbidity (severe underlying disease and/or immunodeficiency)	B	IIht	[37,41,63,66]	Comorbidity: wide variety of risk factors described/investigated, including cancer, cognitive impairment, cardiovascular, respiratory and kidney disease [41]. Chronic pulmonary disease, chronic renal disease and diabetes mellitus [66]. History of malignancy [37]. Previous operative therapy, inflammatory bowel disease and intravenous immunoglobulin treatment [63].

<sup>a</sup>SoR: degree of recommendation to use a (clinical) characteristic as a prognostic marker.

**TABLE 6. Prognostic markers that can be used to determine (increased risk of) recurrent *Clostridium difficile* infection (CDI)**

Characteristics	SoR <sup>a</sup>	QoE	Ref (s) not exhaustive	Comment(s)
Age ( $> 65$ years)	A	IIrh	[42,43,46,67]	Meta-analysis: [43]. Systematic review: [46]. Prospective validation study of risk factor: [42].
Continued use of (non-CDI) antibiotics after diagnosis of CDI and/or after CDI treatment	A	IIrh	[42,43]	Meta-analysis: [43]. Prospective validation study of risk factor: [42].
Comorbidity (severe underlying disease) and/or renal failure	A	IIh	[42,45,68]	Prospective validation study of risk factor: comorbidity conditions rated by Horns' index (scoring system for underlying disease severity) [42].
A history of previous CDI (more than one recurrence)	A	IIt	[26,40,69–71]	Data from randomized controlled trials: [26,70]. Meta-analysis of pivotal randomized controlled trials [40].
Concomitant use of antacid medications (proton pump inhibitors)	B	IIrh	[43,72]	Meta-analysis on recurrent CDI: [43]. Meta-analysis on CDI: [72].
Initial disease severity	B	IIth	[42,67]	Prospective validation study of risk factor [42]. Long-term population based cohort study [67].

<sup>a</sup>SoR: degree of recommendation to use a (clinical) characteristic as a prognostic marker.

colitis mentioned in Table 4 is present, and/or when one or more unfavourable prognostic factors (Table 5) is present:

1. Marked leucocytosis (leucocyte count  $> 15 \times 10^9/L$ )
2. Decreased blood albumin ( $< 30$  g/L)
3. Rise in serum creatinine level ( $\geq 133$   $\mu M$  or  $\geq 1.5$  times the premorbid level)

*Clostridium difficile* infection without signs of severe colitis in older patients ( $\geq 65$  years), serious comorbidity, ICU admission, or immunodeficiency may also be regarded as increased risks of developing severe CDI.

## Treatment of *Clostridium difficile* Infection

Once CDI is diagnosed in a patient, immediate implementation of appropriate infection control measures is mandatory to prevent further spread within the hospital. These include early diagnosis of CDI, surveillance, education of staff, appropriate use of isolation precautions, hand hygiene, protective clothing, environmental cleaning and cleaning of medical equipment, good antibiotic stewardship, and specific measures during

outbreaks. Measures for the prevention and control of CDI ('bundle approach') have been described in an ESCMID guideline by Vonberg et al. [73].

Additional treatment measures include [1,3,4,72,74]:

- Discontinuation of unnecessary antimicrobial therapy
- Adequate replacement of fluid and electrolytes
- Avoidance of anti-motility medications
- Reviewing proton pump inhibitor use

In general it is difficult to compare studies on the treatment of CDI because of the use of variable diagnostic criteria, patient selection and subgroup definitions, stringency of searches for potential enteropathogens, severity of CDI, comorbidities, exposures to causative or concomitant antibiotics, and follow up. Moreover, studies have employed different definitions of clinical and/or microbiological cure and recurrence [2,75]. The variability in definitions and criteria of randomized controlled trials of antibiotic therapy for CDI is illustrated in Table 7. In 13/17 randomized controlled trials of antibiotic treatment of initial CDI, recurrences and duration of follow up were defined. Follow up varied from 3 to 6 weeks



**TABLE 7. Randomized controlled trials of antibiotic treatment of initial *Clostridium difficile* infection (CDI): definitions and criteria of recurrences, follow up and severity of infection**

Trial	Recurrences before study	Relapse/recurrences and follow up	Severity of CDI	Severe CDI excluded/included
[76]	Previous PMC excluded	Recurrences not defined and follow up not specified	Not defined	Not specified
[77]	Not described	Reappearance of diarrhoea <21 days	Not defined	Not specified
[78]	Not described	Reappearance of diarrhoea <5 weeks	Not defined	Not specified
[79]	Not described	Reappearance of diarrhoea after therapy	Not defined	Not specified
[80]	Not described	Follow up: length not clear 'Recurrence of disease': not further specified	No definition but judged by physician	Severe/moderate CDI included, mild CDI excluded
[81]	Not described	Follow up not defined	Not defined	Not specified
[82]	Not described	Not described No follow-up period	Not defined	Not specified
[83]	Treatment for CDI <6 weeks excluded	Reappearance of diarrhoea and other symptoms $\geq$ 1 month	Not defined	Not specified
[84]	Not described	Follow up not further specified Cure followed by return of inclusion criteria CDI <4 weeks	Not defined	Not specified
[85]	Not described	Reappearance of diarrhoea and other symptoms <25–30 days	Severity estimated by: number/shape stool, CRP, WBC, ESR	Severe and mild CDI included. Results for PMC specified
[86]	CDI $\leq$ 6 months excluded	Reappearance of diarrhoea during 28–33 days	Not defined	Not specified. Severe 'medical conditions' excluded Toxic megacolon excluded
[87]	Not specified Excluded oral vanco/metro treatment <7 days before study (at least two doses included)	Reappearance of symptoms <31 days after start of treatment and after at least one negative CD toxin test before retreatment	Not defined	
[88]	Previous CDI excluded	Recurrence of diarrhoea during 30 days	Not defined	Not specified. Ileus and toxic megacolon excluded
[89]	Prior failure of treatment for CDI with study drugs excluded	Recurrence of CD toxin-positive diarrhoea within 21 days	Severe CDI defined as severity assessment score $\geq$ 2 (points). Based on: age (1), temperature (1), Alb (1), WBC (1), endoscopic PMC (2), ICU (2)	Severe and mild CDI included: results specified Life-threatening abdominal complications excluded
[90]	More than one recurrence or relapse within 3 months before study excluded	Recurrence of CD toxin-positive diarrhoea <6 weeks	Severity CDI based on: stools/day, vomiting, ileus, severe abdominal tenderness, WBC, toxic megacolon, life-threatening CDI	Mild to moderately severe CDI included: results not specified Very severe CDI excluded
[91]	More than one recurrence <3 months before study excluded Results specified for CDI <90 days before study.	Return of symptoms (toxin-positive diarrhoea) <31 days after onset of treatment, or clinical response after empiric re-treatment	Severe CDI defined as severity assessment score $\geq$ 2 (points). Based on: age (1), stools/day (1), temperature (1), Alb (1), WBC (1)	Severe and mild CDI included: results specified Unstable vital signs or ICU excluded.
[70]	More than one CDI <3 months before study excluded. Results specified for patients with/without CDI <3 months before study.	Reappearance of CD toxin-positive diarrhoea <4 weeks and need for retreatment for CDI	Mild, moderate and severe CDI: based on bowel movements/day, WBC	Mild, moderate and severe disease included: results specified. Life-threatening or fulminant CDI and toxic megacolon excluded
[91]	More than one CDI <3 months before study excluded Results specified for patients with CDI <3 months before study.	Return of CD toxin-positive diarrhoea <30 days and need for retreatment for CDI	Severe and not-severe CDI based on ESCMID criteria [1]: WBC, creatinine, temperature	Severe and not-severe disease included: results specified for severity. Life-threatening or fulminant CDI and toxic megacolon excluded

Alb, serum albumin; CRP, C-reactive protein; ESR, erythrocyte sedimentation rate; ICU, intensive care unit; PMC, pseudomembranous colitis; WBC, white blood cell count.

after treatment for CDI. In 6/17 randomized controlled trials definitions for severity of disease were given. In most of the studies very severe and/or life-threatening CDI was excluded.

A Cochrane analysis published in 2011 reviewed 15 studies on the antibiotic treatment for CDI in adults [2]. The risk of bias was rated high in 12 of the 15 included studies. The authors concluded that a specific recommendation for the antibiotic treatment of CDI could not be made. Nevertheless, and in spite of the observed limitations, it is apparent that a clear and up-to-date guideline on the treatment of CDI is urgently needed for clinical practice. For this purpose the strength of a recommendation and the quality of evidence are assigned in two separate evaluations in this guideline, hence allowing an assessment of the strength of a recommendation independent of the level of supportive evidence (Tables 1 and 2).

To improve clinical guidance in the treatment of CDI, treatment recommendations are specified for various patient groups:

- A. Initial CDI: non-severe disease
- B. Severe CDI
- C. First recurrence or (risk of) recurrent CDI
- D. Multiple recurrent CDI
- E. Treatment of CDI when oral administration is not possible

The following treatment options are considered:

1. Oral and non-oral antibiotics
2. Toxin-binding resins and polymers
3. Immunotherapy
4. Probiotics
5. Faecal or bacterial intestinal transplantation



## A. Initial *Clostridium difficile* Infection : non-severe Disease

### Oral antibiotic therapy for non-severe disease

**Evidence.** The antibiotics commonly used to treat CDI are oral metronidazole or oral vancomycin.

Oral metronidazole has been shown to be effective in inducing a clinical response and has the advantage of low cost and is assumed to be associated with reduced vancomycin-resistant enterococci (VRE) selection risk. In a pooled intention-to-treat analysis (treating exclusions, deaths and relapses as treatment failures) of three randomized controlled trials comparing symptomatic cure between metronidazole and vancomycin [77,84,88], no statistically significant differences were found [2,75]. Symptomatic cure was achieved in 79% of patients who received vancomycin compared with 71% of patients who received metronidazole (three studies; 335 patients; RR 0.91; 95% CI 0.81–1.03,  $p$  0.14) [2]. However, a recently presented pooled analysis of data from two phase three randomized controlled trials on the use of tolevamer, comparing resolution of diarrhoea and abdominal pain (clinical success) for vancomycin versus metronidazole, showed that overall metronidazole was inferior to vancomycin [92]. Vancomycin significantly improved clinical success (81.1% vs 72.7%; OR 1.681; 95% CI 1.114–2.537;  $p$  0.0134). In addition a retrospective analysis of case records of hospitalized patients with CDI showed that the symptomatic response time was significantly ( $p < 0.01$ ) shorter in patients treated with vancomycin (3.0 days,  $n = 22$ ) compared with those given metronidazole (4.6 days,  $n = 28$ ) [23]. Oral metronidazole is usually recommended for treatment of non-severe disease, whereas oral vancomycin is generally preferred for treatment of severe infections [1,3–5].

Decreased clinical effectiveness of metronidazole treatment for specific ribotypes causing CDI, e.g. PCR ribotype 027, has been described [93]. Although changes in antibiotic resistance and ribotype prevalence have been reported, *in vitro* studies indicate that MICs of metronidazole and vancomycin for endemic *C. difficile* have remained relatively low over the years. Brazier *et al.* concluded that the MICs of metronidazole and vancomycin were not indicative of clinical failure, but MICs for epidemic ribotypes (027, 106 and 001) were several dilutions higher [94]. Indeed there is increasing evidence of the emergence of reduced susceptibility to metronidazole in some *C. difficile* strains, with evidence for clonal spread [95]. Notably, MIC methodology is crucial to the detection of reduced susceptibility to metronidazole; E-tests in particular underestimate the MIC [95,96]. There is also evidence of inferior microbiological efficacy of metroni-

dazole in comparison with vancomycin [21,22]. Although poor gut concentrations of metronidazole alongside reduced susceptibility to metronidazole could explain reduced treatment efficacy, treatment failures have not been associated with decreased susceptibility [95,97,98]. A case–control study found no significant differences in clinical outcome for CDI cases from which strains with reduced susceptibility to metronidazole were recovered versus matched (metronidazole-susceptible) controls [99]. Response to metronidazole was generally poor (slow and prone to recurrence) and the frail elderly patients had a 21% 30-day mortality. However, much larger study groups are needed to determine the clinical significance of CD isolates with reduced susceptibility to metronidazole [99].

Orally administered vancomycin is poorly absorbed from the gastrointestinal tract, and therefore luminal drug levels are high and orders of magnitude are greater than the susceptibility breakpoint concentration for all strains of *C. difficile* tested so far, thereby resulting in a more rapid suppression of *C. difficile* to undetectable levels during therapy and faster resolution of diarrhoea [22,23]. Metronidazole, on the other hand, is well absorbed from the gastrointestinal tract. Mean antibiotic concentrations reported in faeces of patients receiving oral metronidazole range from  $<0.25$  to 9.5 mg/L, and drug concentrations in faeces decrease to undetectable levels as mucosal inflammation improves and diarrhoea resolves [100]. Increased MIC for metronidazole could therefore have implications on clinical cure or recurrences in CDI. Although there are no published reports in which treatment failure has been linked to antimicrobial metronidazole resistance in *C. difficile*, the pharmacokinetic properties of vancomycin are considered superior to those of metronidazole in severe *C. difficile* disease [88].

There is concern that use of vancomycin may be more likely to promote colonization and transmission of VRE by selection pressure. However, both oral metronidazole and oral vancomycin have been associated with the promotion of persistent overgrowth of VRE in stool samples obtained from colonized patients during CDI treatment, thereby increasing the risk of transmission [101]. In a small study of VRE-colonized patients with CDI, who experienced frequent faecal incontinence, skin and environmental VRE contamination was common during and after resolution of diarrhoea. It was concluded that the frequency of VRE contamination of skin or the environment was similar between patients treated with metronidazole ( $n = 17$ ) and those given vancomycin ( $n = 17$ ), although the study clearly had only limited power to examine this issue [102]. In a large retrospective analysis, increased vancomycin use during an outbreak of CDI was not associated with an increase in VRE colonization during a follow-up period of 2 years after the

outbreak period [103]. The authors concluded that restriction of vancomycin use during CDI outbreaks because of the fear of increasing VRE colonization might not be warranted. However, the interpretation of the data was complicated by an outbreak of VRE (VanA) cases that was observed after approximately 20 months of increasing preferential use of vancomycin. As the rate of VanA cases subsequently decreased very quickly, the investigators concluded that this temporary increase reflected a localized clonal outbreak unrelated to the CDI therapy at that time [103].

Although vancomycin and metronidazole are effective in the treatment of CDI, they are both broader-spectrum agents that cause significant disruption of the commensal colonic microbiota. A disruption in the commensal microbiota may predispose to recurrent CDI and intestinal colonization by health-care-associated pathogens such as VRE and *Candida* species. Fidaxomicin appears to cause less disruption of the anaerobic colonization microbiota, and has activity against many VRE strains [104] so it is suggested that the risk of colonization with and transmission of VRE associated with fidaxomicin treatment may be lower compared with vancomycin therapy. A recent study concluded that fidaxomicin was indeed less likely than vancomycin to promote acquisition of VRE and *Candida* species during CDI treatment. However, selection of pre-existing subpopulations of VRE with elevated fidaxomicin MICs was more common during fidaxomicin therapy [105].

Similar cure rates have been demonstrated for oral vancomycin and oral teicoplanin [82,84]. For bacteriological cure, oral teicoplanin may even be more effective than vancomycin [2,82]. Both glycopeptides are active *in vitro* against *C. difficile* isolates [106]. Since 2013 teicoplanin does have a licensed indication for CDI and is available for oral administration. Teicoplanin is not available in the USA. For the purpose of this treatment guideline only oral vancomycin is included in the treatment recommendations.

Tables 8 and 9 report the evidence for oral treatment of initial CDI from randomized trials and observational studies with comments on methodology.

Although oral metronidazole absorption is very high and potentially can lead to more systemic side-effects, adverse effects of oral metronidazole are commonly mild to moderate in severity. The most common adverse reactions reported involve the gastrointestinal tract [107]. Rarely, particularly in association with long duration therapy, metronidazole has been linked to more severe safety issues, e.g. peripheral and optic neuropathy [108] and interactions with warfarins [109].

Oral vancomycin has been shown to be poorly absorbed in most patients, usually producing minimal or subtherapeutic serum concentrations. However, bowel inflammation may

enhance absorption of oral vancomycin, particularly in those with renal failure, thereby increasing the risk for systemic side-effects [110]. A recently performed safety analysis of fidaxomicin in comparison with oral vancomycin revealed no differences in serious adverse events between these agents [111]. Fidaxomicin is minimally absorbed. While no specific concerns related to hypersensitivity reactions were identified during the drug development, hypersensitivity reactions associated with fidaxomicin use have been reported to the FDA in the post-marketing phase. The fidaxomicin labeling was revised to include information about the possibility of hypersensitivity reactions [112].

To evaluate the clinical outcomes of the main antimicrobial agents used in the treatment of CDI, we compared dosages, cure rate, recurrence rate, stated time to response and adverse events of treatment with vancomycin, metronidazole and fidaxomicin. Only randomized controlled trials of antibiotic treatment of initial CDI were included. Results are summarized in Table 10.

**Recommendations.** In case of *non-severe CDI* (no signs of severe colitis) in *non-epidemic situations* and with *CDI clearly induced by the use of antibiotics*, it may be acceptable to stop the inducing antibiotic and observe the clinical response for 48 h, but patients must be followed very closely for any signs of clinical deterioration and placed on therapy immediately if this occurs. Metronidazole is recommended as oral antibiotic treatment of initial CDI in mild/moderate disease. For detailed recommendations on oral antibiotic treatment of initial non-severe CDI refer to Table 11.

#### **Alternative treatment regimens treatment for non-severe disease**

**Evidence.** Tables 12 and 13 report the evidence from randomized trials and observational studies on the non-antibiotic treatment of initial CDI, with comments on methodology. The majority of these alternative treatment strategies are combined with antibiotic treatment.

Currently there are no randomized controlled trials on the use of human intravenous gammaglobulins (IVIG). Passive immunizations with IVIG have been reported to be successful in small case series, but the grade of evidence and strength of recommendation of IVIG are too weak to allow recommendations on the use of IVIG in CDI [4,130]. Hypogammaglobulinaemia, e.g. following solid organ transplants, may predispose to CDI. For this subgroup of patients, IVIG may be beneficial, but more studies are needed before this can be recommended definitively [4].

A recent systematic review on the use of probiotics suggests that probiotics are associated with a reduction in

**TABLE 8.** Randomized controlled trials of oral antibiotic treatment of initial *Clostridium difficile* infection (CDI). Initial cure rate and sustained response rates as a percentage of all patients and relapse rate as a percentage of initially cured patients

Trial	Treatment	Number of patients	Cure (%)	Recurrence (%)	Sustained response (%)
[76]	Vancomycin, 125 mg four times daily, 5 days Placebo	9 7	78 14	0 –	78 –
	No clear case definition. No description of allocation of treatment. Only data of patients with toxin-positive stool shown. Unclear length of follow up and incidence or relapse in placebo group. $p < 0.02$ for comparison of cure rates.				
[77]	Vancomycin, 500 mg four times daily, 10 days Metronidazole 250 mg four times daily, 10 days	32 32	100 97	19 6	81 91
	Only data of patients with toxin-positive stools or pseudomembranous colitis shown. Per-protocol analysis. Follow up 21 days. Differences not statistically significant.				
[78]	Vancomycin, 125 mg four times daily, 7 days Bacitracin, 20 000 U four times daily, 7 days	21 21	86 76	33 42	58 44
	Double-blind. 25% drop-out during follow up of bacitracin group. Follow up 5 weeks. Differences not statistically significant.				
[79]	Vancomycin, 500 mg four times daily, 10 days Bacitracin, 25 000 U four times daily, 10 days	15 15	100 80	20 42	80 46
	Double-blind. Patients had leucocytosis, fever or abdominal pain. 29% drop-out in vancomycin group, 12% in bacitracin group. Per-protocol analysis. Unclear definition of failure ('worsening during treatment'). Failing patients crossed over to alternate drug. Interruption of study drug in vancomycin group for a mean of 2.8 days and in bacitracin group for a mean of 1.8 days. Unclear length of follow up. Differences not statistically significant.				
[80]	Vancomycin, 125 mg four times daily, mean 10.6 days Vancomycin, 500 mg four times daily, mean 10.1 days	24 22	100 100	21 18	79 82
	Variable duration of therapy. 18% dropout rate. Per-protocol analysis. Unclear length of follow up. Differences not statistically significant.				
[81]	Vancomycin, 500 mg twice daily, 10 days Rifaximin, 200 mg three times daily, 10 days	10 10	100 90	– –	– –
	Article in Italian. Patients had diarrhoea, abdominal pain and fever. No description of allocation of treatment. Unclear definition of cure. Differences not statistically significant.				
[82]	Vancomycin, 500 mg four times daily, 10 days Teicoplanin, 100 mg twice daily, 10 days	20 26	100 96	20 8	80 88
	No description of allocation of treatment. Per-protocol analysis. Unclear length of follow up ('at least 1 month'). Differences not statistically significant.				
[83]	Teicoplanin, 100 mg four times daily, 3 days, followed by 100 mg twice daily, 4 days Teicoplanin, 100 mg twice daily, 7 days	24 23	96 70	35 50	62 35
	Double-blind. Outcome of 'improvement, but not cure' (two loose stools per day or one loose stool per day with fever or cramps) was counted as failure. Three patients with improvement in twice daily group; one in four times daily group. Follow up 5 weeks. $p 0.08$ for comparison of cure rates.				
[84]	Vancomycin, 500 mg three times daily, 10 days Metronidazole, 500 mg three times daily, 10 days Teicoplanin, 400 mg twice daily, 10 days Fusidic acid, 500 mg three times daily, 10 days	31 31 28 29	94 94 96 93	17 17 7 30	78 78 89 65
	Follow up 30 days. Only statistically significant difference was relapse rate of fusidic acid versus teicoplanin ( $p 0.042$ ).				
[85]	Metronidazole, 400 mg three times daily, 7 days Fusidic acid, 250 mg three times daily, 7 days	55 59	93 83	30 30	65 58
	Double-blind. 13% drop-out during treatment; 15% further drop-out during follow up. Per-protocol analysis. Follow up 35 days. Differences not statistically significant.				
[86]	Metronidazole, 250 mg four times daily, 10 days Nitazoxanide, 500 mg twice daily, 7 days Nitazoxanide, 500 mg twice daily, 10 days	34 40 36	82 90 89	30 6 16	57 67 75
	No definition of relapse. Double-blind. 23% drop-out during treatment. Per-protocol analysis. Follow up 31 days. Differences not statistically significant.				
[87]	Metronidazole, 500 mg three times daily, 10 days Metronidazole, 500 mg three times daily + rifampicin 300 mg twice daily, 10 days	20 19	65 63	38 42	40 37
	Intention-to-treat analysis. Follow up 40 days. Differences not statistically significant.				
[88]	Vancomycin, 125 mg four times daily, 10 days Metronidazole, 250 mg four times daily, 10 days	71 79	97 84	7 14	90 72
	Double-blind. 13% drop-out during treatment. Per-protocol analysis. Follow up 21 days. $p 0.006$ for comparison of cure rates. $p 0.27$ for comparison of relapse rates. The original protocol was stratified in a group with mild and a group with severe disease (based on age, fever, albumin level and leucocyte count), which resulted in a larger difference between cure rates in the group with severe disease and a statistically non-significant difference between cure rates in the group with mild disease. Intention-to-treat analysis with dropouts regarded as failures resulted in a statistically significant difference between overall cure rates (initial cure minus relapse; 57 out of 90 versus 64 out of 82; risk ratio 0.91). Other comparisons were not significant anymore in the intention-to-treat analysis.				
[89]	Fidaxomicin, 50 mg twice daily, 10 days Fidaxomicin, 100 mg twice daily, 10 days Fidaxomicin, 200 mg twice daily, 10 days	14 15 16	71 80 94	8 0 6	65 80 88
	Open-label. Patients with signs of highly severe CDI (>12 bowel movements per day, vomiting, severe abdominal tenderness, ileus, white blood cell count >30, toxic megacolon) were excluded. Cure = complete resolution of diarrhoea. Follow up 6 weeks after end of treatment.				
[90]	Vancomycin, 125 mg four times daily, 10 days Nitazoxanide, 500 mg twice daily, 10 days	27 22	74 77	7 5	69 73
	CDI = stool EIA for toxin A or B positive AND (temperature >38.3°C OR abdominal pain OR leucocytosis). Patients with more than one episode in preceding 6 months were excluded. 12% dropout rate during treatment. Double-blind, placebo-controlled. Modified intention-to-treat analysis. Industry-sponsored. Cure = complete resolution of symptoms during 3 days after completion of therapy. Per-protocol analysis: 87 versus 94% cure. Follow up 31 days after start of treatment. No differences in severity subgroups. Differences not statistically significant.				
[70]	Vancomycin, 125 mg four times daily, 10 days Fidaxomicin, 200 mg twice daily, 10 days	309 287	86 88	25 15	65 75
	Placebo-controlled. Industry-sponsored. Very severe CDI and more than one previous episode excluded. Designed as non-inferiority trial. 4 weeks follow up for recurrences after completion of study drug. Cure = <4 times daily passage of unformed stools AND no necessity for additional treatment. Fidaxomicin was not associated with fewer recurrences in CDI due to PCR ribotype 027 as opposed to non-027. Modified intention-to-treat (patients who received at least one dose of the study drug) and per-protocol analyses were similar.				
[91]	Vancomycin, 125 mg four times daily, 10 days Fidaxomicin, 200 mg twice daily, 10 days	257 252	87 88	27 13	64 77
	Methods identical to the trial by Louie et al. [70]. Contrary to that trial, this trial did show fewer recurrences in both polymerase chain reaction ribotype 027 and non-027 patients, although the difference was not significant for the former subgroup.				

antibiotic-associated diarrhoea [131]. A recent meta-analysis on probiotic prophylaxis for CDI, concluded that moderate-quality evidence suggests a beneficial effect of probiotic prophylaxis in CDI without an increase in clinically important

adverse events [132]. However, a Cochrane analysis concluded that there was insufficient evidence to recommend probiotics, in general, as an adjunct to antibiotics in the treatment of *C. difficile* diarrhoea [133]. Although no cases of

**TABLE 9. Observational studies of oral antibiotic treatment of initial *Clostridium difficile* infection (CDI). Initial cure rate and sustained response as a percentage of all patients and relapse rate as a percentage of initially cured patients**

Trial	Treatment	Number of patients	Cure (%)	Recurrence (%)	Sustained response (%)
Antibiotics					
[113]	Vancomycin	79	96	14	83
[114]	Vancomycin	16	100	13	87
[115]	Metronidazole	13	100	15	85
[116]	Vancomycin	189	97	24	74
[106]	Vancomycin 500 mg four times daily, 10 days	23	100	13	87
	Teicoplanin 200 mg twice daily, 10 days	22	100	0	100
[117]	Metronidazole	632	98	6	92
	Vancomycin	122	99	10	89
[57]	Metronidazole	44	?	50	–
[118]	Metronidazole	99	62	?	–
[119]	Metronidazole	207	78	28	56
[68]	Metronidazole	1123	84	29	60
	Vancomycin	112	?	28	–
[120]	Fidaxomicin varying dose	45	91	5	86
[121]	Nitazoxanide 500 mg twice daily, 10 days	35	74	27	54
	Patients first failed metronidazole				
[101]	Metronidazole	34	>90	12	>79
	Ten patients switched to vancomycin				
	Vancomycin	18	>90	11	>80
[122]	Tigecycline varying duration	4	100	0	100
	Severe CDI. Follow up at least 3 months				
[123]	Rifaximin 400 mg three times daily 2 weeks follow up	8	100	10	90

translocation of microorganisms have been reported in clinical trials with probiotics for antibiotic-associated diarrhoea or CDI, probiotics should be used with caution. Several studies of invasive disease have been reported, resulting from the use of probiotics such as *Saccharomyces boulardii* in debilitated or immunocompromised patients [134,135]. Moreover, probiotics were associated with increased mortality, partly due to non-occlusive mesenteric ischaemia, in a randomized controlled trial in acute pancreatitis [136].

**Recommendations.** There is insufficient evidence to support administration of probiotics, toxin-binding resins and polymers, or monoclonal antibodies. For detailed recommendations refer to Table 14.

## B: Severe *Clostridium difficile* Infection

### Oral antibiotic therapy

**Evidence.** In 6/17 randomized controlled trials, severity of disease was defined. Definitions varied among the studies.

Only in 4/6 of these trials were treatment results specified for severity of disease (Table 15).

**Recommendations.** Based on its pharmacokinetic properties vancomycin is considered superior to metronidazole in severe *C. difficile* disease [22,88]. The use of high doses of vancomycin (500 mg orally four times daily) was included in the Infectious Diseases Society of America/Society for Healthcare Epidemiology of America treatment guidelines [3] for management of severe complicated CDI as defined by the treating physician. However, there is insufficient evidence to support the use of doses >125 mg four times daily in the absence of ileus [80].

Fidaxomicin was not inferior to vancomycin for initial cure of CDI, but there are no data available on the efficacy of this drug in severe life-threatening disease [70,91].

For detailed recommendations on oral antibiotic treatment of severe CDI refer to Table 16.

### Surgery for complicated *Clostridium difficile* infection

**Evidence.** Patients with fulminant CDI who fail to respond and who progress to systemic toxicity, peritonitis, or toxic colonic dilatation and bowel perforation require surgical intervention [4]. Mortality rates of emergency surgery in complicated CDI remain high, ranging from 19% to 71% depending on the clinical condition of the patient at the time of surgery [138]. However, recently a systemic review of the existing literature was performed to assess the effect on mortality of colectomy for the treatment of fulminant CDI. The authors concluded that colectomy is associated with a lower mortality than continued medical treatment when this is no longer improving the patient [139]. Several studies suggest that earlier colectomy (time from presentation to surgery) is associated with improved survival [140]. Independent risk factors for mortality in patients who underwent colectomy that have been found among multiple studies include: the development of shock (need for vasopressors), increased serum lactate ( $\geq 5$  mM), mental status changes, end organ failure, renal failure and the need for preoperative intubation and ventilation [29,35,138,141,142]. The more negative prognostic signs a patient has, the earlier surgical consultation and operative management should be considered. The established operative management of severe, complicated CDI has been subtotal colectomy with end-ileostomy [140]. However, recently an alternative surgical treatment with creation of a diverting loop ileostomy, followed by colonic lavage, has been shown to reduce morbidity and mortality, while preserving the colon. The surgical approach involves the laparoscopic creation of a diverting loop ileostomy. The colon is then lavaged in an ante-grade fashion through the ileostomy with a high volume

**TABLE 10.** Results of randomized controlled trials of oral antibiotic treatment of initial *Clostridium difficile* infection (CDI) with vancomycin/teicoplanin, metronidazole and fidaxomicin: comparison of dosages, cure rate, recurrence rate, stated time to response or adverse effects due to treatment

	Trial	Number of patients	Dosages and duration of therapy	Time to initial response (mean)	Cure rate (%)	Recurrence rate (%) and definition	Adverse events (%)
Vancomycin	[76]	9	125 mg four times daily 5 days	–	78	0 Recurrence not defined, follow-up period not specified	–
	[77]	32	500 mg four times daily 10 days	3.2 days	100	19 Reappearance of diarrhoea <21 days after therapy	3 Drug intolerance
	[78]	21	125 mg four times daily 7 days	–	86	33 Reappearance of diarrhoea <5 weeks after therapy	–
	[79]	15	500 mg four times daily 10 days	–	100	20 Reappearance of diarrhoea after therapy	–
	[80]	24	125 mg four times daily mean 11 days	4 days	100	21 Follow-up: length not clear	0
		22	500 mg four times daily mean 10 days	4 days	100	18 Recurrence of disease not further specified	0
	[81]	10	500 mg twice daily 10 days	3.8 days	100	Follow up not defined Not described	0
	[82]	20	500 mg four times daily 10 days	3.6 days	100	4 No follow-up period	0
	[84]	31	500 mg three times daily 10 days	3.1 days	94	17 Reappearance of diarrhoea and other symptoms >= 1 month after therapy. Follow up not further specified	0
	[88]	71	125 mg four times daily 10 days	–	97	7 Recurrence of CD toxin-positive diarrhoea within 21 days after start of therapy	1 (nausea)
	[90]	27	125 four times daily 10 days	Median: 96 h	74	7 Return of symptoms (toxin-positive diarrhoea) <31 days after onset of treatment, or clinical response after empiric re-treatment for CDI	0
	[70]	30	125 mg four times daily 10 days	Median: 78 h	86	25 Reappearance of CD toxin-positive diarrhoea <4 weeks after treatment and need for retreatment for CDI	Possibly or definitely related: nine serious events related to laboratory test results: 1.2
[91]	257	125 mg four times daily 10 days	Median: 58 h	87	27 Return of CD toxin positive diarrhoea <30 days after treatment and need for retreatment for CDI	Any treatment-emergent adverse event related to study drug: 13.8	
Teicoplanin	[82]	26	100 mg twice daily 10 days	3.4 days	96	2 Reappearance of diarrhoea and other symptoms >= 1 month after therapy. Follow up not further specified	0
	[84]	28	400 mg twice daily 10 days	2.8 days	96	7 Reappearance of diarrhoea and other symptoms < 25–30 days after therapy	0
	[83]	24	100 mg four times daily, 3 days, followed by 100 mg twice daily, 4 days	–	96	35	7–8 vomiting, nausea, exanthema, arthralgia, pruritus, hallucinations. No abnormal laboratory results
Metronidazole		23	100 mg twice daily 7 days	–	70	50	
	[77]	32	250 mg four times daily 10 days	3.1 days	97	6 Reappearance of diarrhoea <21 days after therapy	3
	[84]	31	500 mg three times daily 10 days	3.2 days	94	17 Reappearance of diarrhoea and other symptoms <25–30 days after therapy	10 Gastrointestinal discomfort
[85]	55	400 mg three times daily 7 days	Within 5 days	93	30 Reappearance diarrhoea	14.5 Gastrointestinal discomfort, exanthema, taste	



Table 10 (Continued)

Trial	Number of patients	Dosages and duration of therapy	Time to initial response (mean)	Cure rate (%)	Recurrence rate (%) and definition	Adverse events (%)	
[86]	34	250 mg twice daily 10 days	Median: 3 days	82	during 28–33 days after treatment 30 Reappearance of symptoms <31 days after start of treatment and after at least one negative CD toxin test before retreatment	Related to study drug: 0 serious adverse events not related to study drug: 18.2 intolerance or allergy: 0	
[87]	20	500 mg three times daily 10 days	6.6 days	65	38 Recurrence of diarrhoea <30 days after treatment	40 (not specified if related to study drug: rash, nausea vomiting)	
[88]	79	250 mg four times daily 10 days	–	84	14 Recurrence of CD toxin-positive diarrhoea <21 days after start of therapy	1.3 (nausea)	
Fidaxomicin	[89]	14	50 mg twice daily 10 days	Median 6.3 days	71	8	20 but not related to study drug
		15	100 mg twice daily 10 days	Median 4.8 days	80	0	
		16	200 mg twice daily 10 days	Median 3.6 days	94	6 Recurrence of CD toxin-positive diarrhoea <6 weeks after treatment	
[70]	287	200 mg twice daily 10 days	Median 58 h in the MITT	88	15 Reappearance of CD toxin-positive diarrhoea <4 weeks and need for retreatment for CDI	Possibly or definitely related: 9.7 Serious events related to laboratory test results: 4.7	
[91]	252	200 mg twice daily 10 days	Median 56 h	88	13 Return of CD toxin-positive diarrhoea <30 days and need for retreatment for CDI	Any treatment-emergent adverse event related to study drug: 11.7	

TABLE 11. Recommendations on oral antibiotic treatment of initial *Clostridium difficile* infection (CDI): non-severe disease

Treatment	SoR	QoE	Ref(s)	Comment(s)
Metronidazole, 500 mg three times daily 10 days	A	I	[77,84–88]	No statistically significant difference in cure rate between metronidazole and vancomycin or teicoplanin. Statistically significant difference in sustained clinical cure between metronidazole and vancomycin in favour of vancomycin in one study [2,88] (and pooled results of two randomized controlled trials published only in abstract form [92,123,124]).
Vancomycin, 125 mg four times daily 10 days	B	I	[70,76,78,80,82,84,88,90,91]	Cochrane analysis: teicoplanin significantly better than vancomycin for bacteriological cure and borderline superior in terms of symptomatic cure [2].
Fidaxomicin, 200 mg twice daily 10 days	B	I	[70,89,91]	Evidence limited to two Phase III studies. Fewer recurrences as compared to vancomycin, except for <i>C. difficile</i> PCR ribotype 027 [91].
Vancomycin, 500 mg four times daily 10 days	C	I	[77,79–82,84]	Vancomycin: Equal cure rate 500 mg four times daily orally compared with 125 mg four times daily orally [80].
Stop inducing antibiotic(s) and observe the clinical response for 48 h	C	II	[116,117]	Rate of spontaneous resolution unknown in mild CDI. Studies performed before increased incidence of hypervirulent strains.

of polyethylene glycol 3350 or balanced electrolyte solution and the effluent is collected via a rectal drainage tube. A catheter is placed in the efferent limb of the ileostomy to deliver vancomycin flushes in an antegrade fashion in the postoperative period. In addition, patients receive intravenous metronidazole for 10 days [143]. A multicentre randomized controlled trial is currently being conducted to provide level I evidence for possible implementation of this new treatment into standard practice [<http://clinicaltrials.gov/show/NCT01441271>].

**Recommendations.** Total abdominal colectomy should be performed to treat CDI in case of:

- Perforation of the colon
- Systemic inflammation and deteriorating clinical condition despite maximal antibiotic therapy; this includes the clinical diagnoses of toxic megacolon, acute abdomen and severe ileus. Colectomy should preferably be performed before colitis becomes very severe. Serum lactate may, *inter alia*, serve as a marker for severity (operate before lactate exceeds 5.0 mM).

A future alternative to colectomy may be diverting loop ileostomy and colonic lavage, combined with antibiotic treatment (intracolonic antegrade vancomycin and intravenous metronidazole).

**TABLE 12. Randomized controlled trials of alternative treatment regimens for initial *Clostridium difficile* infection (CDI). Initial cure rate and sustained response as a percentage of all patients and relapse rate as a percentage of initially cured patients**

Trial	Treatment	Number of patients	Cure (%)	Recurrence (%)	Sustained response (%)
Probiotics [126]	Vancomycin or metronidazole + <i>Saccharomyces boulardii</i> 2 × 10 <sup>10</sup> CFU/day, 4 weeks	31	–	19	–
	Vancomycin or metronidazole + placebo	33	–	24	–
Double-blind. No control for type, duration or dose of antibiotic. Unclear definition of relapse. Follow up 8 weeks after start of treatment. p 0.86 for comparison of relapse rates.					
Toxin-binding resins and polymers:					
[24]	Tolevamer 1 g three times daily, 14 days + placebo	94	60	16	50
	Tolevamer 2 g three times daily, 14 days + placebo	91	79	7	74
	Vancomycin 125 mg four times daily, 10 days + placebo	94	91	19	74
Non-inferiority trial. Patients with stool frequency >12 daily or abdominal pain were excluded. Tolevamer could be prolonged when inciting antibiotic could not be stopped. Double-blind. 23% drop-out. Per-protocol analysis. Cure rate of tolevamer 2 g non-inferior in comparison with vancomycin (Chow-test p 0.03). Non-inferiority of tolevamer 1 g compared with vancomycin could not be demonstrated. p 0.05 for comparison of relapse rates of tolevamer 2 g with vancomycin. Relapse rates of tolevamer 1 g and vancomycin not statistically different. Follow up 6–8 weeks.					
[124] <sup>a</sup>	Tolevamer, 3 g three times daily, 14 days	266	47	3	46
	Vancomycin, 125 mg four times daily, 10 days	134	81	23	62
[125] <sup>a</sup>	Metronidazole, 375 mg four times daily, 10 days	143	72	27	53
	Tolevamer, 3 g three times daily, 14 days	268	42	6	40
	Vancomycin, 125 mg four times daily, 10 days	125	81	18	66
Immunotherapy [71]	Metronidazole, 375 mg four times daily, 10 days	135	73	19	59
	Single dose of 10 mg/kg CDA1 and CDB1 (intravenously administered human monoclonal antibodies against TcdA and TcdB) with standard antimicrobial therapy	101	93	7	87
	Placebo with standard antimicrobial therapy	99	87	25	65
Industry-sponsored and -analysed. Patients must have diarrhoea and receive vancomycin or metronidazole at time of enrolment. Diarrhoea at least two unformed stools on two consecutive days or more than six unformed stools on 1 day. Recurrence = new episode of diarrhoea with new positive stool toxin test after resolution of initial diarrhoea. Analysis for recurrence only performed in those who were cured, received >7 days of antimicrobial therapy and did not receive intravenous immunoglobulin (93 versus 82). Dropout rate 9 versus 13%, mainly due to deaths not related to CDI. Only 30% (n = 30) of patients treated with vancomycin received monoclonal antibodies versus 22% (n = 22) placebo. Follow up 12 weeks. p <0.001 for comparison of relapse rates. Intention-to-treat analysis. Primary endpoint was changed during the study before unblinding. Original endpoint: resolution of illness. Subgroup analysis: similar results, although difference much smaller in inpatients than outpatients. Length of hospitalization did not differ.					

<sup>a</sup>Poster presentation.

**TABLE 13. Observational studies of alternative treatment regimens for initial *Clostridium difficile* infection (CDI). Initial cure rate as a percentage of all patients and relapse rate as a percentage of initially cured patients**

Trial	Treatment	Number of patients	Cure (%)	Recurrence (%)
Toxin-binding resins and polymers [127]	Colestipol 10 g four times daily, 5 days	12	25	–
	Originally set up as a randomized placebo-controlled trial. Placebo group was merged with historical control, however. Only six patients had toxin-positive stool.			
Passive immunotherapy with immune whey: [128]	Metronidazole or vancomycin followed by immune whey protein concentrate, 14 days	16	100	0
	56% of patients had recurrent CDI; mean follow up 333 days.			
[129]	Metronidazole or vancomycin followed by immune whey protein concentrate, 14 days	109	100	10
	109 episodes; 101 patients; 40% of patients had recurrent CDI.			

### C: First Recurrence or (Risk of) recurrent *Clostridium difficile* Infection

#### Oral antibiotic therapy

**Evidence.** In 3/17 randomized controlled trials of antibiotic treatment of initial CDI, results were specified for CDI before the study (Table 17).

**Recommendations.** The incidence of a second recurrence after treatment of a first recurrence with oral metronidazole or vancomycin is similar. Fewer secondary recurrences

with oral fidaxomicin as compared with vancomycin after treatment of a first recurrence are reported [70,91,144]. However, the evidence on fidaxomicin for this specific subgroup of CDI patients is limited to two phase III studies and based on a retrospective subset analysis of data and a limited number of patients (number of patients in the modified intention-to-treat analysis: fidaxomicin n = 79 and vancomycin n = 80) [144]. There are no prospective randomized controlled trials performed with metronidazole, vancomycin or fidaxomicin in this specific patient group. In addition, fidaxomicin was not associated with fewer recur-



**TABLE 14. Recommendations on alternative treatment regimens for initial *Clostridium difficile* infection (CDI)**

Type of intervention	Treatment	SoR	QoE	Ref(s)	Comment(s)
Immunotherapy	Human monoclonal antibodies against TcdA and TcdB with standard oral antimicrobial therapy (metronidazole and vancomycin)	C	I	[71]	Evidence limited to Phase II randomized controlled trial. Primary endpoint changed during study. Reduced recurrence of CDI: analysis for recurrence only performed in those who were cured, received >7 days of antimicrobial therapy and did not receive intravenous gammaglobulins
	Passive immunotherapy with immune whey after standard oral antimicrobial therapy	C	II	[129]	Observational study: 101 CDI patients (40% recurrent CDI). Results suggest reduction in recurrence rate.
Probiotics	Oral vancomycin or oral metronidazole + <i>Saccharomyces boulardii</i>	D	I	[126,137]	Comparison of relapse rates: in subgroup analysis efficacy in recurrent CDI, but not in initial CDI. Evidence-based review: [137].
Toxin-binding resins and polymers	Tolevamer, 3 g three times daily	D	I	[24]	Evidence limited to Phase II randomized controlled trial. Non-inferiority study: tolevamer versus vancomycin.

**TABLE 15. Randomized controlled trials of oral antibiotic treatment of initial *Clostridium difficile* infection (CDI) in which severity of disease is defined and outcome of treatment is specified for severity of diseases**

Study	Treatment	CDI severity: moderate/mild (M), severe (S) Nr of patients (%)	Initial cure No. of patients (%)	Relapse No. of patients (% of patients with initial cure)	Sustained response rate <sup>a</sup> No. of patients (% of all patients)
[88]	Vancomycin, 125 mg four times daily, 10 days	M 40/71 (56)	39/40 (98)	2/39 (5)	37/40 (93)
	Metronidazole 250 mg four times daily, 10 days	S 31/71 (44)	30/31 (97)	3/30 (10)	27/31 (87)
		M 41/79 (52)	37/41 (90)	3/37 (8)	34/41 (83)
		S 38/79 (48)	29/38 (76)	6/29 (21)	23/38 (61)
Intention-to-treat analysis:	Vancomycin 125 mg four times daily, 10 days	M 44/82 (49)	39/44 (89)	2/39 (5)	37/44 (84)
	Metronidazole 250 mg four times daily, 10 days	S 38/82 (46)	30/38 (79)	3/30 (10)	27/38 (71)
		M 46/90 (51)	37/46 (80)	3/37 (8)	34/46 (74)
		S 44/90 (49)	29/44 (66)	6/29 (21)	23/44 (52)
[90]	Vancomycin, 125 mg four times daily, 10 days	M 17/27 (63)	13/17 (76)	1/13 (8)	12/17 (71)
	Nitazoxanide 500 mg twice daily, 10 days	S 10/27 (37)	7/10 (70)	1/7 (14)	6/10 (60)
		M 12/22 (55)	9/12 (75)	0/9 (0)	9/12 (75)
[70]	Vancomycin 125 mg four times daily, 10 days	S 10/22 (45)	8/10 (80)	1/8 (13)	7/10 (70)
	Fidaxomicin 200 mg twice daily, 10 days	M 186/309 (60)	156/186 (85)	38/156 (24)	118/186 (63)
		S 123/309 (40)	109/123 (89)	29/109 (27)	80/123 (65)
[91]	Vancomycin 125 mg four times daily, 10 days	M 175/287 (61)	161/175 (92)	27/161 (17)	134/175 (77)
	Fidaxomicin 200 mg twice daily, 10 days	S 112/287 (39)	92/112 (82)	12/92 (13)	80/112 (71)
		M 196/257 (76)	180/196 (92)	46/180 (26)	134/196 (68)
	S 61/257 (24)	43/61 (71)	14/43 (33)	29/61 (48)	
	M 189/252 (75)	173/189 (92)	24/173 (14)	149/189 (79)	
	S 63/252 (25)	48/63 (76)	4/48 (8)	44/63 (70)	

<sup>a</sup>Sustained response rate: clinical cure and no recurrences during follow up.

**TABLE 16. Recommendations on oral antibiotic treatment of initial *Clostridium difficile* infection (CDI): severe disease**

Treatment	SoR	QoE	Ref(s)	Comment(s)
Vancomycin, 125 mg four times daily for 10 days	A	I	[70, 88, 90, 91]	Cure rate higher as compared with metronidazole in severe CDI [88] <sup>a</sup>
Vancomycin 500 mg four times daily for 10 days	B	III (I <sup>b</sup> )	[80]	Randomized controlled trial on dose effectiveness: no significant differences in measurable responses of high-dose compared to low-dose regimens. However: results not stratified for severity of illness [80] <sup>a</sup> .
Fidaxomicin 200 mg twice daily for 10 days	B	I	[70,89,91]	Evidence limited to two Phase III studies [70,91]. Fewer recurrences compared with vancomycin 125 mg four times daily in severe disease (except for PCR ribotype 027). No data on the efficacy in severe life-threatening disease and/or toxic megacolon: excluded from both studies.
Metronidazole, 500 mg three times daily for 10 days	D	I	[88]	Cure rate lower as compared with vancomycin in severe CDI [88]. Intention to treat analysis not reported. Extremely severe CDI excluded <sup>a</sup> . Differences in symptomatic cure of metronidazole versus vancomycin not statistically significant in a pooled analysis [2]. ICU admission and hypoalbuminaemia (= disease severity) predictors of metronidazole failure [119].

<sup>a</sup>Two studies reported in abstract form confirm the superiority of vancomycin over metronidazole for treatment of (severe) CDI [92,124,125].

rences in CDI due to PCR ribotype 027 as opposed to non-027 in one of the randomized controlled trials [70]. Therefore, based on the evidence currently available, the Strength of Recommendation for treating a first recurrence of CDI with oral vancomycin or oral fidaxomicin is

considered equal (B-I), unless disease has progressed from non-severe to severe.

For detailed recommendations on oral antibiotic treatment of mild/moderate initial CDI with risk for recurrent CDI or a first recurrence refer to Table 18.

## D: Multiple recurrent *Clostridium difficile* Infection

### Antibiotic and non-antibiotic treatment strategies

**Evidence.** Tables 19 and 20 report the evidence from randomized trials and observational studies with comments on methodology.

**Recommendations.** In non-severe second (or later) recurrences of CDI oral vancomycin or fidaxomicin is recommended. Vancomycin and fidaxomicin are equally effective in resolving CDI symptoms, but fidaxomicin has been shown to be associated with a lower likelihood of CDI recurrence after a first recurrence [104,144]. However, there are no prospective randomized controlled trials investigating the efficacy of fidaxomicin in patients with *multiple* recurrences of CDI. Vancomycin is preferably administered using a tapered and/or pulsed regimen.

Recently the first randomized controlled trial on faecal enteric instillation has been published: faecal transplantation following antibiotic treatment with an oral glycopeptide is reported to be highly effective in treating multiple recurrent CDI [145].

For detailed recommendations on treatment regimens of multiple recurrent CDI refer to Tables 21 and 22.

## E: Treatment of *Clostridium difficile* Infection when oral Administration is not possible

### Evidence

Metronidazole remains the only parenteral antibiotic therapy supported by case series [192]. Intravenous metronidazole (500 mg intravenous three times daily) may be added to oral vancomycin, if the patient has ileus or significant abdominal distension [4,44]. However, there are no randomized controlled trials available to guide this recommendation.

It is still unknown how to best treat patients with ileus due to CDI. There are some anecdotal reports on delivery of vancomycin to the gut by means other than orally, mainly through intracolonic delivery. Questions regarding the efficacy, optimal dosing and duration of treatment with intracolonic vancomycin remain unanswered [193,194]. Prospective clinical trials with other antibiotics, like tigecycline, have not yet been performed to support general use [122,195].

**TABLE 17.** Randomized controlled trials of antibiotic treatment of initial *Clostridium difficile* infection (CDI) in which relapses are defined and outcome of treatment is specified for CDI before study

Study	Treatment	CDI before study, No. of patients (%)	Initial cure No. of patients (%)	Relapse No. of patients (% with initial cure)	Sustained response rate <sup>a</sup> No. of patients (%)
[90]	Vancomycin, 125 mg four times daily, 10 days	5/27 (19)	4/5 (80)	1/4 (25)	3/5 (60)
	Nitazoxanide, 500 mg twice daily, 10 days	2/22 (9)	2/2 (100)	1/2 (50)	1/2 (50)
[70]	Vancomycin, 125 mg four times daily, 10 days	54/309 (17)	48/54 (89)	15/48 (31)	33/54 (61)
	Fidaxomicin 200 mg twice daily, 10 days	48/287 (17)	42/48 (88)	9/42 (21)	33/42 (78)
[91]	Vancomycin 125 mg four times daily, 10 days	36/257 (14)	32/36 (89)	11/32 (34)	21/36 (58)
	Fidaxomicin 200 mg twice daily, 10 days analysed in: [144]	40/252 (16)	37/40 (93)	7/37 (19)	30/40 (75)

<sup>a</sup>Sustained response rate: clinical cure and no recurrences during follow up.

**TABLE 18.** Recommendations on oral antibiotic treatment of mild/moderate initial *Clostridium difficile* infection (CDI) with risk for recurrent CDI or first recurrence

Treatment	SoR	QoE	Ref(s)	Comment(s)
Vancomycin, 125 mg four times daily for 10 days	B	I	[70,82,90,91]	No statistically significant difference in recurrence rate between vancomycin and teicoplanin [2,82,84].
Fidaxomicin, 200 mg twice daily for 10 days	B	I	[70,89,91]	Evidence limited to two Phase III studies [70,91]. Retrospective subset analysis: fewer secondary recurrences with fidaxomicin ( $n = 16/79$ patients) as compared with vancomycin ( $n = 26/80$ patients) after treatment of a first recurrence [144]. Fidaxomicin was not associated with fewer recurrences in CDI due to PCR ribotype 027 as opposed to non-027 [70].
Metronidazole, 500 mg three times daily for 10 days	C	I	[27,88]	Recurrence rate: metronidazole not inferior to vancomycin for treatment of mild primary CDI [2,82,88] or after a first recurrence [27]. Vancomycin significantly more effective in bacteriological cure than metronidazole in recurrent CDI [69].
Vancomycin, 500 mg four times daily for 10 days	C	III	[80]	One randomized controlled trial on dose effectiveness in primary CDI: no significant differences in responses of high-dose compared with low-dose regimens vancomycin. However, results not stratified for recurrent CDI [80].

**TABLE 19. Randomized controlled studies of treatment of recurrent *Clostridium difficile* infection (CDI)**

Trial	Treatment	No. of patients	Failure <sup>a</sup> [%]
<b>Faecal or bacterial instillation</b>			
[145]	Vancomycin 500 mg four times daily, 14 days	13	69
	Vancomycin 500 mg four times daily 14 days + bowel lavage	13	77
	Vancomycin 500 mg four times daily 4 days + bowel lavage + nasoduodenal infusion donor faeces	16	19
	3/16 patients with failure after first donor faeces infusion received second infusion from a different donor: 2/3 resolved. Treatment with donor faeces was superior to either of the vancomycin regimens (both $p < 0.001$ ). Open label. No definition of diarrhoea. Study terminated by use of Haybittle-Peto rule at unplanned interim analysis. Fecotherapy group was older, had more comorbidities, higher creatinine, and more infections with PCR ribotype 027. Other characteristics were comparable.		
<b>Probiotics</b>			
[126]	Vancomycin or metronidazole + <i>Saccharomyces boulardii</i> $2 \times 10^{10}$ CFU/day, 4 weeks	26	35
	Vancomycin or metronidazole + placebo	34	65
	Double-blind. No control for type, duration or dose of antibiotic. Unclear definition of relapse. Follow up 8 weeks after start of treatment. $p = 0.04$ for comparison of failure rates.		
[146]	Vancomycin 500 mg four times daily, 10 days, followed by <i>Saccharomyces boulardii</i> $2 \times 10^{10}$ CFU/day, 4 weeks	18	17
	Vancomycin 500 mg four times daily, 10 days, followed by placebo	14	50
	Vancomycin 125 mg four times daily, 10 days, followed by <i>Saccharomyces boulardii</i> $2 \times 10^{10}$ CFU/day, 4 weeks	45	51
	Vancomycin 125 mg four times daily, 10 days, followed by placebo	38	45
	Metronidazole 1 g/day, 10 days, followed by <i>Saccharomyces boulardii</i> $2 \times 10^{10}$ CFU/day, 4 weeks	27	48
	Metronidazole 1 g/day, 10 days, followed by placebo	26	50
	Follow up 5 months after completion of study. $p = 0.05$ for the comparison of failure rates in patients who received 500 mg vancomycin four times daily. 22% drop-out in this group. No further statistically significant differences.		
[147]	Metronidazole 400 mg three times daily, 10 days + <i>Lactobacillus plantarum</i> 299v $5 \times 10^{10}$ CFU/day, 38 days	12	42
	Metronidazole 400 mg three times daily, 10 days + placebo	9	67
	Double-blind. 28% drop-out. Follow up 70 days. Difference not statistically significant.		
[148]	Vancomycin or metronidazole followed by <i>Lactobacillus GG</i> $6 \times 10^{11}$ CFU/day, 21 days	8	38
	Vancomycin or metronidazole followed by placebo	7	14
	Patients blinded. No control for type, duration or dose of antibiotic. Follow up 60 days after completion of antibiotic. Difference not statistically significant.		
<b>Passive immunotherapy with immune whey</b>			
[149]	Colostrum immune whey 200 mL three times daily + placebo, 14 days	18	44
	Metronidazole 400 mg three times daily + placebo, 14 days	20	45
	Double-blind. Multi-centre trial. Follow up 70 days. Difference not statistically significant.		

<sup>a</sup>Non-response or relapse.**TABLE 20. Observational studies for treatment of recurrent *Clostridium difficile* infection (CDI)**

Trial	Treatment	No. of patients	Failure <sup>b</sup> (%)	Mean follow up
<b>Antibiotics</b>				
[150]	Vancomycin taper, 21 days, followed by vancomycin pulse, 21 days	22	0	6 months
[151]	Vancomycin 125 mg four times daily + rifampicin 600 mg twice daily, 7 days	7	0	12 months
[69]	Vancomycin 1–2 g/day	14	71	59 days
	Vancomycin <1 g/day	48	54	59 days
	Vancomycin $\geq 2$ g/day	21	43	59 days
	Vancomycin taper	29	31	80 days
	Vancomycin pulse	7	14	80 days
	Metronidazole <1 g/day	29	45	59 days
	Metronidazole 1.5 g/day	5	40	59 days
	Metronidazole 2 g/day	2	0	59 days
[152]	Vancomycin, 14 days, followed by rifaximin varying dose, 14 days	8	13	233 days
[153]	Rifaximin 400 mg three times daily, 14 days, followed by rifaximin 200 mg three times daily, 14 days	5	0	310 days
	Rifaximin 400 mg three times daily, 36 days	1	100	–
[154]	Rifaximin 400 mg three times daily, 14 days	25	36	56 days
	Severe CDI excluded. Patients unresponsive to metronidazole 500 mg three times daily, 5 days. Cure = negative stool PCR for TcdB. All patients had resolution of diarrhoea, but no definition or description of how this was measured is given.			
<b>Probiotics</b>				
[155]	Metronidazole or bacitracin, 10 days, followed by <i>Lactobacillus GG</i> $10^{10}$ CFU/day, 7–10 days	5	20	–
[156]	<i>Lactobacillus GG</i> $6 \times 10^8$ CFU/day, 14 days	4	0	11 months
<b>Faecal or bacterial instillation<sup>a</sup></b>				
[157]	Faecal enema	16	19	(5 days–3 years)
	faecal enema $n = 15$ , enteric tube $n = 1$			
[158]	Faecal or bacterial enema	6	0	6 months
	two faecal and four bacterial mixture			

Table 20 (Continued)

Trial	Treatment	No. of patients	Failure <sup>b</sup> (%)	Mean follow up
[159]	Rectal tube	7	0	2 year
[160]	Faecal instillation through colonoscope or gastrostoma	18	17	–
[161]	Lower gastrointestinal tract	6	0	(9–50 months)
[162]	Nasogastric tube, median three courses two patients died: not CDI-related, 15/16 cure after first faecal transplantation (FT), 1 relapse	16	6	90 days
[163]	Faecal enema	5	0	–
[164] <sup>b</sup>	Rectal catheter	45	4	(≤1 year)
[165]	Colonoscopy, enema Complete resolution of symptoms in 8/16 and marked reduction in 7/16	16	6	6 week
[166]	Vancomycin 500 mg four times daily, followed by faecal instillation by nasoduodenal tube or colonoscopy	7	29 0 after repeated infusion	150 days
[167]	Nasogastric tube	12	17	90 days
[164] <sup>c</sup>	Faecal enema CDI in refractory inflammatory bowel disease (IBD)	6	0	8 week
[168]	Nasogastric tube	15	27	Median 4 months
[169]	Colonoscopy	37	8	12 months
[170]	Colonoscopy 1/19 non-responders after first FT; all cured after second FT	19	5	27 months
[171]	Enema	7	0	9 months
[172]	Colonoscopy	13	15	5 months
[173]	Colonoscopy	12	0	(3 week–8 year)
[174]	Gastroscopy or colonoscopy	40	27	80 days
[175]	Colonoscopy	26	8	11 months
[176]	Colonoscopy 7/77 treatment failures within 90 days after treatment (early recurrence). 8/77 recurrence >90 days after treatment (late recurrence).	77	19	17 months
[177]	Faecal enema	27	7	427 days
[178]	5/27 patients had two FT: 2/5 failures Faecal instillation through coloscope Patients with (14) and without (29) IBD. 6/43 patients had two FT: 2/6 failures	43	14	2 months
[179]	Colonoscopy Initial failures were all PCR-ribotype 027.	70	11	1 year
<b>Immunotherapy</b>				
[180]	Intravenous gammaglobulin 400 mg/kg every 3 weeks, 4–6 months	5	0	5 months
[181]	Intravenous gammaglobulin 400 mg/kg day 1 and 21 Intravenous gammaglobulin, varying dose	4 5	0 40	7.5 months 2.8 months
[56]	Intravenous gammaglobulin 300–500 mg/kg, 1–6 doses	5	40	86 days
[182]	Intravenous gammaglobulin 150–400 mg/kg once	14	71	6.6 months
[183]	Intravenous gammaglobulin 200–300 mg/kg once	18	33 (died or colectomy)	–
[184]	Intravenous gammaglobulin 75–400 mg/kg, 1–5 days	21	57 (died)	–

Non-response or relapse.  
<sup>a</sup>Reviewed by Refs. [164,185–191].  
<sup>b</sup>Louie (2008) abstract only derived from Ref. [164].  
<sup>c</sup>Borody (2008) abstract only derived from Ref. [164].

TABLE 21. Recommendations on oral antibiotic treatment of multiple recurrent *Clostridium difficile* infection (CDI) (more than one relapse)

Treatment	SoR	QoE	Ref(s)	Comment(s)
Vancomycin, 125 mg four times daily for 10 days, followed by pulse regimen (125–500 mg/day every 2–3 days) for at least 3 weeks.	B	IIt	[69,150]	Retrospective case cohort of two placebo/antibiotic trials [69]: [126,146]. Observational study: [150]. Expert opinion [3].
Vancomycin, 125 mg four times daily for 10 days, followed by taper regimen: gradually decreasing the dose to 125 mg per day.	B	IIt	[69,150]	Retrospective case cohort of two placebo/antibiotic trials [69]: [125,146]. Observational study: [150]. Expert opinion [3].
Fidaxomicin, 200 mg twice daily for 10 days	B	IIrt	[75,144]	Evidence limited to two Phase III studies [70,91]. Retrospective subset analysis: fewer recurrences as compared to vancomycin treatment after first recurrence [144]. Systematic review: [75]. Efficacy after multiple recurrences was not investigated [144].
Vancomycin, 500 mg four times daily for 10 days	C	IIrt	[69,75]	Retrospective case cohort of two placebo/antibiotic trials: [126,146]. Trend for lower recurrence frequency for high-dose vancomycin [69]. Systematic review: [75].
Metronidazole, 500 mg three times daily for 10 days	D	IIrt	[69,75]	Retrospective case cohort of two placebo/antibiotic trials: [126,146]. Trend for lower recurrence frequency for high-dose vancomycin and low-dose metronidazole [69]. Systematic review: [75].

**TABLE 22.** Recommendations on non-antibiotic treatment (in combination with antibiotic treatment) of recurrent *Clostridium difficile* infection (CDI) (more than one relapse)

Type of intervention	Treatment	SoR	QoE	Ref(s)	Comment(s)
Faecal or bacterial instillation	Vancomycin, 500 mg four times daily, 4 days + bowel lavage + nasoduodenal infusion donor faeces	A	I	[145]	Also many observational studies and meta-analyses. [164,186,189–191].
Probiotics	Vancomycin or metronidazole + <i>Saccharomyces boulardii</i>	D	I	[126]	Comparison of relapse rates: in subgroup analysis efficacy in recurrent CDI, but <i>not</i> in initial CDI. Evidence-based review: [137].
	Vancomycin or metronidazole + <i>Lactobacillus</i> spp.	D	I	[147,148]	Evidence-based review: [137].
Passive immunotherapy with immune whey	Colostrum immune whey	D	I	[149]	Study interrupted early.

**TABLE 23.** Recommendations on non-oral antibiotic treatment of initial *Clostridium difficile* infection (CDI): mild and severe disease

Patient subgroup	Treatment	SoR	QoE	Ref(s)	Comment(s)
Non-severe disease	Intravenous metronidazole 500 mg three times daily for 10 days	A	IIu	[192]	Retrospective uncontrolled study [192].
Severe disease and/or complicated or refractory CDI	Intravenous metronidazole 500 mg three times daily for 10 days + vancomycin retention enema 500 mg in 100 mL normal saline four times daily intracolonic for 10 days	A	IIru	[192–194]	Retrospective uncontrolled study [192].
		B	III		Systematic review [193,194]. Expert opinion [3].
	Intravenous metronidazole 500 mg three times daily for 10 days + vancomycin 500 mg in 100 mL normal saline four times daily by oral/nasogastric tube for 10 days	A	IIru	[192–194]	Retrospective uncontrolled study [192].
		B	III		Systematic review [193,194]. Expert opinion [3].
Intravenous tigecycline 50 mg twice daily for 14 days	C	III	[122]	Observational study/case report [122].	

### Recommendations

When oral treatment is not possible, parenteral metronidazole is recommended, preferably combined with intracolonic or nasogastric administration of vancomycin. Parenteral tigecycline as salvage therapy is only recommended with marginal strength. For detailed recommendations refer to Table 23.

### Summary of Definitions

**Episode of CDI.** A clinical picture compatible with CDI and microbiological evidence of free toxins and the presence of *C. difficile* in stool, without reasonable evidence of another cause of diarrhoea.

or

Pseudomembranous colitis diagnosed during endoscopy, after colectomy or on autopsy.

*Clinical pictures compatible with CDI.*

**Diarrhoea:** loose stools, i.e. taking the shape of the receptacle or corresponding to Bristol stool chart types 5–7, plus a stool frequency of three stools in 24 or fewer consecutive hours, or more frequently than is normal for the individual.

**Ileus:** signs of severely disturbed bowel function such as vomiting and absence of stool with radiological signs of bowel distension.

**Toxic megacolon:** radiological signs of distension of the colon (>6 cm in transverse width of colon) and signs of a severe systemic inflammatory response.

**Severe CDI.** Severe or life-threatening CDI is defined as an episode of CDI with (one or more specific signs and symptoms of) severe colitis or a complicated course of disease, with significant systemic toxin effects and shock, resulting in need for ICU admission, colectomy or death.

One or more of the following unfavourable prognostic factors can be present without evidence of another cause:

- Marked leucocytosis (leucocyte count  $>15 \times 10^9/L$ )
- Decreased blood albumin ( $<30$  g/L)
- Rise in serum creatinine level ( $\geq 133 \mu M$  or  $\geq 1.5$  times the pre-morbid level)

**Recurrent CDI.** Recurrence is present when CDI re-occurs <8 weeks after the onset of a previous episode, provided the symptoms from the previous episode resolved after completion of initial treatment.

**Treatment response.** Treatment response is present when after therapy either stool frequency decreases or stool consistency improves and parameters of disease severity (clinical, laboratory, radiological) improve and no new signs of severe disease develop.

Treatment response should be observed daily and evaluated after at least 3 days, assuming that the patient is not worsening on treatment. Treatment with metronidazole, in particular, may result in a clinical response only after 3–5 days. After clinical response, it may take weeks for stool consistency and frequency to become entirely normal.

## Summary of Treatment Recommendations

Strength of Evidence (SoE: I to III) and Strength of Recommendation (SoR: A to D) are shown in brackets. For grading definitions we refer to Tables 1 and 2.

### A: Initial *Clostridium difficile* Infection: non-severe Disease

#### Non-antibiotic treatment

In non-epidemic situations and with (non-severe) CDI clearly induced by the use of antibiotics, it may be acceptable to stop the inducing antibiotic and observe the clinical response for 48 h, but patients must be followed very closely for any signs of clinical deterioration and placed on therapy immediately if this occurs. (C-II).

#### Oral antibiotic treatment

Metronidazole orally 500 mg three times daily for 10 days (A-I)

Vancomycin orally 125 mg four times daily for 10 days (B-I)

Fidaxomicin orally 200 mg twice daily for 10 days (B-I)

### B: Severe *Clostridium difficile* Infection

#### Oral antibiotic treatment

Vancomycin orally 125 mg four times daily for 10 days (A-I)

Fidaxomicin orally 200 mg twice daily for 10 days (B-I)

Notes:

- It can be considered to increase the vancomycin dosage to 500 mg four times daily for 10 days (B-III)
- There is no evidence that supports the use of fidaxomicin in life-threatening CDI (D-III)

The use of oral metronidazole in severe CDI or life-threatening disease is strongly discouraged (D-I).

#### Surgical treatment

Total abdominal colectomy with ileostomy should be performed in case of:

- Perforation of the colon
- Systemic inflammation and deteriorating clinical condition not responding to antibiotic therapy; including toxic megacolon, an acute abdomen and severe ileus.

Surgical treatment should preferably be performed before colitis becomes very severe. Serum lactate may, *inter alia*, serve as a marker for severity (operate before lactate exceeds 5.0 mM).

A future alternative to colectomy may be diverting loop ileostomy and colonic lavage, combined with *antibiotic treatment* (intracolonic antegrade vancomycin and intravenous metronidazole).

### C: First Recurrence or (Risk of) recurrent *Clostridium difficile* Infection

#### Oral antibiotic treatment

Fidaxomicin orally 200 mg twice daily for 10 days (B-I)

Vancomycin orally 125 mg four times daily for 10 days (B-I)

Metronidazole orally 500 mg three times daily for 10 days (C-I)

Note: Fidaxomicin was not associated with fewer recurrences in CDI due to PCR ribotype 027 as opposed to non-027 ribotypes.

### D: Multiple recurrent *Clostridium difficile* Infection

#### Oral antibiotic treatment

Fidaxomicin orally 200 mg twice daily for 10 days (B-II)

Vancomycin orally 125 mg four times daily for 10 days followed by pulse strategy (B-II)

or

Vancomycin orally 125 mg four times daily for 10 days followed by taper strategy (B-II)

#### Non-antibiotic treatment in combination with oral antibiotic treatment

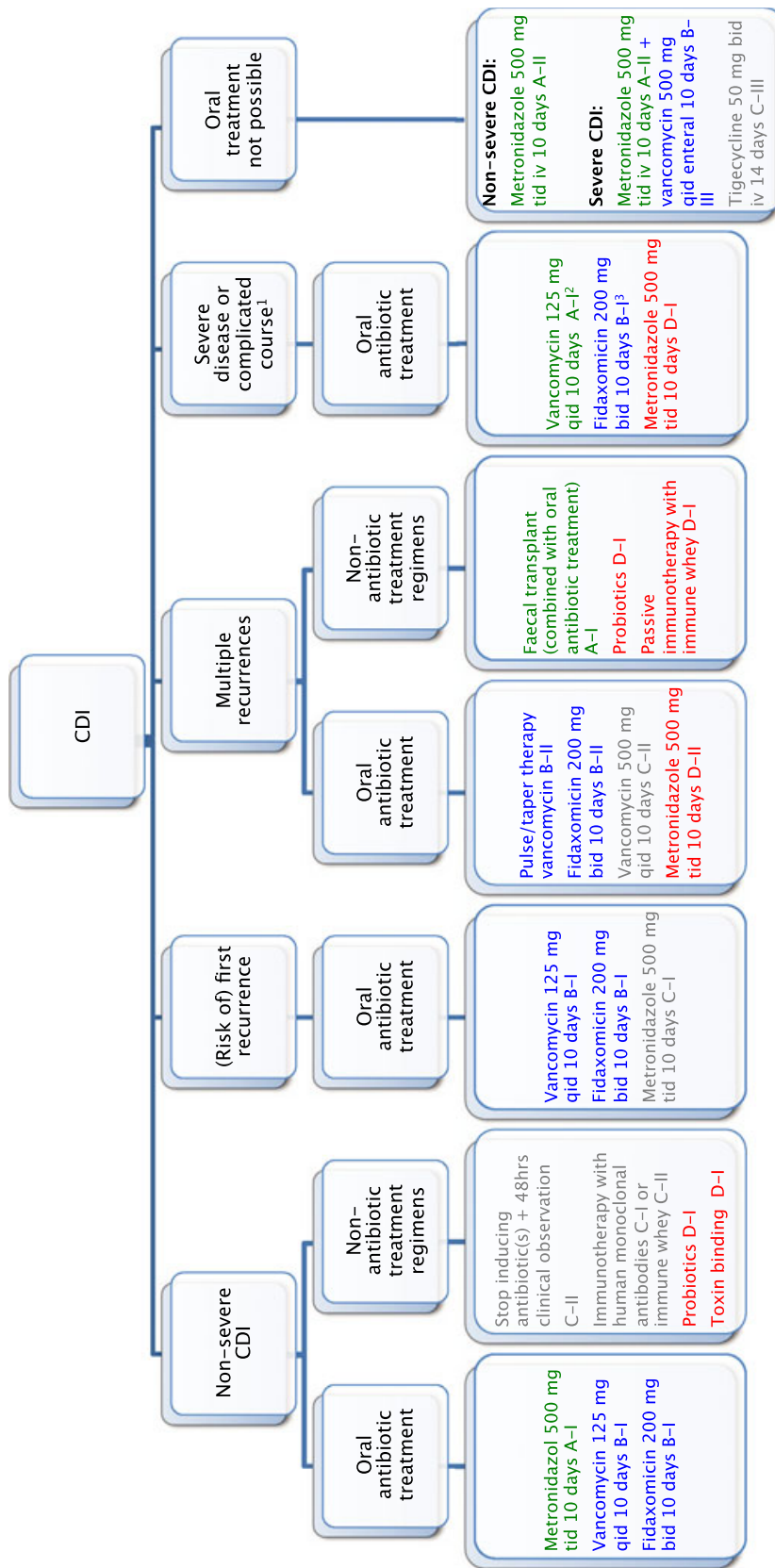
For multiple recurrent CDI unresponsive to repeated antibiotic treatment, faecal transplantation in combination with oral antibiotic treatment is strongly recommended (A-I).

### E: Treatment of *Clostridium difficile* Infection when oral Administration is not possible

#### Antibiotic treatment

Non-severe CDI: intravenous metronidazole 500 mg three times daily for 10 days (A-II).





**FIG. 1.** Schematic overview of therapeutic regimens for *Clostridium difficile* infection (CDI). <sup>1</sup>Severe CDI or complicated course: surgical therapy not included in this overview; <sup>2</sup>It can be considered to increase the oral dosage of vancomycin to 500 mg four times daily for 10 days (B-II); <sup>3</sup>There is no evidence that supports the use of fidaxomicin in life-threatening CDI (D-II); Strength of Recommendation (SoR) A = green (Strongly supports a recommendation for use); SoR B = blue (Moderately supports a recommendation for use); SoR C = grey (Marginally supports a recommendation for use); SoR D=red (Recommendation against use).



Severe CDI: intravenous metronidazole 500 mg three times daily for 10 days (A-II) combined with vancomycin retention enema 500 mg in 100 mL normal saline four times daily intracolonic, or combined with vancomycin 500 mg four times daily by oral/nasogastric tube for 10 days (B-III).

A schematic overview of currently available therapeutic regimens for CDI, including the quality of evidence (QoE: I to III) and strength of recommendations (SoR: A to D) are shown in Fig. 1.

## Authorship

Four draft versions of this guideline document were written by three authors (MB, EK, JvD) and critiqued by the Expert Panel. A consensus was reached, resulting in the final version.

## Transparency Declaration

Authors: The authors declare that they have no conflicts of interest.

Expert Panel: All members of the expert group completed a Conflict of Interest Disclosure Form (COI).

## Appendix I: On behalf of the Committee

### Expert panel composition

F. Allerberger, Austrian Agency for Health and Food Safety (AGES), Vienna, Austria.

E. Bouza, Department of Infectious Diseases, Madrid, Spain.

J. E. Coia, Department of Clinical Microbiology, Glasgow Royal Infirmary, Glasgow, UK.

O. A. Cornely, Department of Internal Medicine, Clinical Trials Centre Cologne, ZKS Köln, BMBF 01KN1106, Cologne Excellence Cluster on Cellular Stress Responses in Aging-Associated Diseases (CECAD), University Hospital of Cologne, German Centre for Infection Research (DZIF), partner site Bonn-Cologne, Germany.

F. Fitzpatrick, Beaumont Hospital and Health Protection Surveillance Centre, Dublin, Ireland.

B. Guery, Department of Infectious Diseases, Lille, France.

M. Wilcox, Department of Microbiology, Old Medical, School Leeds General Infirmary, Leeds Teaching Hospitals & University of Leeds, Leeds, UK.

D. Nathwani, Department of Infectious Diseases Ninewells Hospital & Medical School, Dundee, UK.

T. Norén, Department of Infectious Diseases, Örebro University Hospital, SE 701 85 Örebro Sweden.

B. Olesen, Department of Microbiology, Herlev Hospital, Herlev, Denmark.

E. Rakoczi, Department of Clinical Pharmacology, Infectious Diseases and Allergology, Kenezy County Hospital, Debrecen, Hungary.

T. Welte, Department of Infectious Diseases, Hannover Medical School, Hannover, Germany.

A. F. Widmer, Department of Infectious Diseases, Universitätsspital, Basel, Switzerland.

## References

- Bauer MP, Kuijper EJ, van Dissel JT. European Society of Clinical Microbiology and Infectious Diseases (ESCMID): treatment guidance document for *Clostridium difficile* infection (CDI). *Clin Microbiol Infect* 2009; 15: 1067–1079.
- Nelson RL. Antibiotic treatment for *Clostridium difficile*-associated diarrhea in adults. *Cochrane Database Syst Rev* 2011; CD004610. doi: 10.1002/14651858.CD004610.pub4.
- Cohen SH, Gerding DN, Johnson S et al. Clinical practice guidelines for *Clostridium difficile* infection in adults: 2010 update by the Society for Healthcare Epidemiology of America (SHEA) and the Infectious Diseases Society of America (IDSA). *Infect Control Hosp Epidemiol* 2010; 31: 431–455.
- Surawicz CM, Brandt LJ, Binion DG et al. Guidelines for diagnosis, treatment, and prevention of *Clostridium difficile* infections. *Am J Gastroenterol* 2013; 108: 478–498.
- Cheng AC, Ferguson JK, Richards MJ et al. Australasian Society for Infectious Diseases guidelines for the diagnosis and treatment of *Clostridium difficile* infection. *Med J Aust* 2011; 194: 353–358.
- Guyatt GH, Oxman AD, Vist GE et al. GRADE: an emerging consensus on rating quality of evidence and strength of recommendations. *BMJ* 2008; 26: 924–926.
- Hsu J, Brožek JL, Terracciano L et al. Application of GRADE: making evidence-based recommendations about diagnostic tests in clinical practice guidelines. *Implement Sci* 2011; 6: 62.
- Ullmann AJ, Cornely OA, Donnelly JP et al. ESCMID guideline for the diagnosis and management of *Candida* diseases 2012: developing European guidelines in clinical microbiology and infectious diseases. *Clin Microbiol Infect* 2012; 18: 1–8.
- Brouwers MC, Kho ME, Browman GP et al. AGREE II: advancing guideline development, reporting, and evaluation in health care. *Prev Med* 2010; 51: 421–424.
- Bartlett JG, Gerding DN. Clinical recognition and diagnosis of *Clostridium difficile* infection. *Clin Infect Dis* 2008; 46 (suppl 1): S12–S18.
- Kuijper EJE, Coignard BB, Tüll PP. Emergence of *Clostridium difficile*-associated disease in North America and Europe. *Clin Microbiol Infect* 2006; 12 (suppl 6): 2–18.
- Crobach MJT, Goorhuis A, Kelly CP et al. European Society of Clinical Microbiology and Infectious Diseases (ESCMID): data review and recommendations for diagnosing *Clostridium difficile*-infection (CDI). *Clin Microbiol Infect* 2009; 15: 1053–1066.
- Wilcox MH, Planche T, Fang FC. What is the current role of algorithmic approaches for diagnosis of *Clostridium difficile* infection? *J Clin Microbiol* 2010; 48: 4347–4353.
- Davies KA, Planche TD, Coen P et al. The largest ever study to define a testing algorithm to optimize the laboratory diagnosis of *C. difficile* infection. In: 22nd European Congress of Clinical Microbiology and

- Infectious Diseases (ECCMID); 2012 in London, UK. Abstract LB2817.
15. Planche TD, Davies KA, Coen PG, et al. Differences in outcome according to *Clostridium difficile* testing method: a prospective multicentre diagnostic validation study of *C difficile* infection. *Lancet Infect Dis* 2013; 13: 936–945.
  16. O'Donnell LJ, Virjee J, Heaton KW. Detection of pseudodiarrhoea by simple clinical assessment of intestinal transit rate. *BMJ* 1990; 300: 439–440.
  17. McDonald LC, Coignard B, Dubberke E, Song X, Horan T, Kutty PK. Recommendations for surveillance of *Clostridium difficile*-associated disease. *Infect Control Hosp Epidemiol* 2007; 28: 140–145.
  18. Lewis SJ, Heaton KW. Stool form scale as a useful guide to intestinal transit time. *Scand J Gastroenterol* 1997; 32: 920–924.
  19. Knoop FC, Owens M, Crocker IC. *Clostridium difficile*: clinical disease and diagnosis. *Clin Microbiol Rev* 1993; 6: 251–265.
  20. Moudgal V, Sobel J. *Clostridium difficile* colitis: a review. *Hosp Pract* 2012; 40: 139–148.
  21. Kuijper EJ, Wilcox MH. Editorial commentary: decreased effectiveness of metronidazole for the treatment of *Clostridium difficile* infection? *Clin Infect Dis* 2008; 47: 63–65.
  22. Nassir AI WN, Sethi AK, Nerandzic MM, Bobulsky GS, Jump RLP, Donskey CJ. Comparison of clinical and microbiological response to treatment of *Clostridium difficile*-associated disease with metronidazole and vancomycin. *Clin Infect Dis* 2008; 47: 56–62.
  23. Wilcox MH, Howe R. Diarrhoea caused by *Clostridium difficile*: response time for treatment with metronidazole and vancomycin. *J Antimicrob Chemother* 1995; 36: 673–679.
  24. Louie TJ, Peppe J, Watt CK et al. Tolevamer, a novel nonantibiotic polymer, compared with vancomycin in the treatment of mild to moderately severe *Clostridium difficile*-associated diarrhea. *Clin Infect Dis* 2006; 43: 411–420.
  25. Kelly CP. Can we identify patients at high risk of recurrent *Clostridium difficile* infection? *Clin Microbiol Infect* 2012; 18 (suppl 6): 21–27.
  26. Fekety R, McFarland LV, Surawicz CM, Greenberg RN, Elmer GW, Mulligan ME. Recurrent *Clostridium difficile* diarrhea: characteristics of and risk factors for patients enrolled in a prospective, randomized, double-blinded trial. *Clin Infect Dis* 1997; 24: 324–333.
  27. Pépin J, Routhier S, Gagnon S, Brazeau I. Management and outcomes of a first recurrence of *Clostridium difficile*-associated disease in Quebec, Canada. *Clin Infect Dis* 2006; 42: 758–764.
  28. Figueroa I, Johnson S, Sambol SP, Goldstein EJC, Citron DM, Gerding DN. Relapse versus reinfection: recurrent *Clostridium difficile* infection following treatment with fidaxomicin or vancomycin. *Clin Infect Dis* 2012; 55 (suppl 2): S104–S109.
  29. Sailhamer EA, Carson K, Chang Y et al. Fulminant *Clostridium difficile* colitis: patterns of care and predictors of mortality. *Arch Surg* 2009; 144: 433–439.
  30. Hall JF, Berger D. Outcome of colectomy for *Clostridium difficile* colitis: a plea for early surgical management. *Am J Surg* 2008; 196: 384–388.
  31. Dallal RM, Harbrecht BG, Boujoukas AJ et al. Fulminant *Clostridium difficile*: an underappreciated and increasing cause of death and complications. *Ann Surg* 2002; 235: 363–372.
  32. Henrich TJ, Krakower D, Bitton A, Yokoe DS. Clinical risk factors for severe *Clostridium difficile*-associated disease. *Emerg Infect Dis* 2009; 15: 415–422.
  33. Kelly MCP, LaMont MJT. *Clostridium difficile* infection. *Annu Rev Med* 1998; 49: 375–390.
  34. Rubin MS, Bodenstern LE, Kent KC. Severe *Clostridium difficile* colitis. *Dis Colon Rectum* 1995; 38: 350–354.
  35. Longo WE, Mazuski JE, Virgo KS, Lee P, Bahadursingh AN, Johnson FE. Outcome after colectomy for *Clostridium difficile* colitis. *Dis Colon Rectum* 2004; 47: 1620–1626.
  36. Miller MA, Louie T, Mullane K et al. Derivation and validation of a simple clinical bedside score (ATLAS) for *Clostridium difficile* infection which predicts response to therapy. *BMC Infect Dis* 2013; 13: 148.
  37. Lungulescu OA, Cao W, Gatskevich E, Tlhabano L, Stratidis JG. CSI: a severity index for *Clostridium difficile* infection at the time of admission. *J Hosp Infect* 2011; 79: 151–154.
  38. Morgan OW, Rodrigues B, Elston T et al. Clinical severity of *Clostridium difficile* PCR ribotype 027: a case–case study. *PLoS ONE* 2008; 3: e1812.
  39. Huttunen R, Vuento R, Syrjänen J, Tissari P, Aittoniemi J. Case fatality associated with a hypervirulent strain in patients with culture-positive *Clostridium difficile* infection: a retrospective population-based study. *Int J Infect Dis* 2012; 16: e532–e535.
  40. Crook DW, Walker AS, Kean Y et al. Fidaxomicin versus vancomycin for *Clostridium difficile* infection: meta-analysis of pivotal randomized controlled trials. *Clin Infect Dis* 2012; 55 (suppl 2): S93–S103.
  41. Welfare MR, Welfare MR, Lalayiannis LC et al. Co-morbidities as predictors of mortality in *Clostridium difficile* infection and derivation of the ARC predictive score. *J Hosp Infect* 2011; 79: 359–363.
  42. Hu MY, Katchar K, Kyne L et al. Prospective derivation and validation of a clinical prediction rule for recurrent *Clostridium difficile* infection. *Gastroenterology* 2009; 136: 1206–1214.
  43. Garey KW, Sethi S, Yadav Y, DuPont HL. Meta-analysis to assess risk factors for recurrent *Clostridium difficile* infection. *J Hosp Infect* 2008; 70: 298–304.
  44. Voelker R. Increased *Clostridium difficile* virulence demands new treatment approach. *JAMA* 2010; 26: 2017–2019.
  45. Bauer MP, Hensgens MPM, Miller MA et al. Renal failure and leukocytosis are predictors of a complicated course of *Clostridium difficile* infection if measured on day of diagnosis. *Clin Infect Dis* 2012; 55 (suppl 2): S149–S153.
  46. Abou Chakra CN, Pepin J, Valiquette L. Prediction tools for unfavourable outcomes in *Clostridium difficile* infection: a systematic review. *PLoS ONE* 2012; 7: e30258.
  47. Miller M, Gravel D, Mulvey M et al. Health care-associated *Clostridium difficile* infection in Canada: patient age and infecting strain type are highly predictive of severe outcome and mortality. *Clin Infect Dis* 2010; 50: 194–201.
  48. Walk ST, Micic D, Jain R et al. *Clostridium difficile* ribotype does not predict severe infection. *Clin Infect Dis* 2012; 55: 1661–1668.
  49. Walker AS, Eyre DW, Crook DW, Peto TE, Wilcox MH. Response to Walk et al. *Clostridium difficile* ribotype does not predict severe infection. *Clin Infect Dis* 2013; 56: 1589–1600.
  50. Walker AS, Eyre DW, Wyllie DH et al. Relationship between bacterial strain type, host biomarkers and mortality in *Clostridium difficile* infection. *Clin Infect Dis* 2013; 56: 1589–1600.
  51. Goorhuis A, Bakker D, Corver J et al. Emergence of *Clostridium difficile* infection due to a new hypervirulent strain, polymerase chain reaction ribotype 078. *Clin Infect Dis* 2008; 47: 1162–1170.
  52. Kelly CP, Kyne L. The host immune response to *Clostridium difficile*. *J Med Microbiol* 2011; 60: 1070–1079.
  53. Sun X, Wang H, Zhang Y, Chen K, Davis B, Feng H. Mouse relapse model of *Clostridium difficile* infection. *Infect Immun* 2011; 79: 2856–2864.
  54. Wullt M, Noren T, Ljungh A, Akerlund T. IgG antibody response to toxins A and B in patients with *Clostridium difficile* infection. *Clin Vaccine Immunol* 2012; 19: 1552–1554.
  55. Wilcox M, Minton J. Role of antibody response in outcome of antibiotic-associated diarrhoea. *Lancet* 2001; 357: 158–159.
  56. Wilcox MH. Descriptive study of intravenous immunoglobulin for the treatment of recurrent *Clostridium difficile* diarrhoea. *J Antimicrob Chemother* 2004; 53: 882–884.

57. Kyne L, Warny M, Qamar A, Kelly CP. Association between antibody response to toxin A and protection against recurrent *Clostridium difficile* diarrhoea. *Lancet* 2001; 357: 189–193.
58. Kyne L, Warny M, Qamar A, Kelly CP. Asymptomatic carriage of *Clostridium difficile* and serum levels of IgG antibody against toxin A. *N Engl J Med* 2000; 342: 390–397.
59. Warny M, Vaerman JP, Avesani V, Delmée M. Human antibody response to *Clostridium difficile* toxin A in relation to clinical course of infection. *Infect Immun* 1994; 62: 384–389.
60. Aronsson B, Granstrom M, Mollby R, Nord CE. Serum antibody response to *Clostridium difficile* toxins in patients with *Clostridium difficile* diarrhoea. *Infection* 1985; 13: 97–101.
61. Leav BA, Blair B, Leney M et al. Serum anti-toxin B antibody correlates with protection from recurrent *Clostridium difficile* infection (CDI). *Vaccine* 2010; 28: 965–969.
62. Mulligan ME, Miller SD, McFarland LV, Fung HC, Kwok RY. Elevated levels of serum immunoglobulins in asymptomatic carriers of *Clostridium difficile*. *Clin Infect Dis* 1993; 16 (suppl 4): S239–S244.
63. Greenstein AJ, Byrn JC, Zhang LP, Swedish KA, Jahn AE, Divino CM. Risk factors for the development of fulminant *Clostridium difficile* colitis. *Surgery* 2008; 143: 623–629.
64. Wanahita A, Goldsmith EA, Musher DM. Conditions associated with leukocytosis in a tertiary care hospital, with particular attention to the role of infection caused by *Clostridium difficile*. *Clin Infect Dis* 2002; 34: 1585.
65. Ramaswamy R, Grover H, Corpuz M, Daniels P, Pitchumoni CS. Prognostic criteria in *Clostridium difficile* colitis. *Am J Gastroenterol* 1996; 91: 460–464.
66. Wenisch JM, Schmid D, Kuo HW et al. Hospital-acquired *Clostridium difficile* infection: determinants for severe disease. *Eur J Clin Microbiol Infect Dis* 2012; 31: 1923–1930.
67. Eyre DW, Walker AS, Wyllie D et al. Predictors of first recurrence of *Clostridium difficile* infection: implications for initial management. *Clin Infect Dis* 2012; 55 (suppl 2): S77–S87.
68. Pépin J, Alary ME, Valiquette L et al. Increasing risk of relapse after treatment of *Clostridium difficile* colitis in Quebec, Canada. *Clin Infect Dis* 2005; 40: 1591–1597.
69. McFarland LV, Elmer GW, Surawicz CM. Breaking the cycle: treatment strategies for 163 cases of recurrent *Clostridium difficile* disease. *Am J Gastroenterol* 2002 Jun 25; 97: 1769–1775.
70. Louie TJ, Miller MA, Mullane KM et al. Fidaxomicin versus vancomycin for *Clostridium difficile* infection. *N Engl J Med* 2011; 364: 422–431.
71. Lowy I, Molrine DC, Leav BA et al. Treatment with monoclonal antibodies against *Clostridium difficile* toxins. *N Engl J Med* 2010; 362: 197–205.
72. Janarthanan S, Ditah I, Adler DG, Ehrinpreis MN. *Clostridium difficile*-associated diarrhea and proton pump inhibitor therapy: a meta-analysis. *Am J Gastroenterol* 2012; 107: 1001–1010.
73. Vonberg RP, Kuijper EJ, Wilcox MH et al. Infection control measures to limit the spread of *Clostridium difficile*. *Clin Microbiol Infect* 2008; 14 (suppl 5): 2–20.
74. Martinez FJ, Leffler DA, Kelly CP. *Clostridium difficile* outbreaks: prevention and treatment strategies. *Risk Manag Healthc Policy* 2012; 5: 55–64.
75. Drekonja DM, Butler M, MacDonald R et al. Comparative effectiveness of *Clostridium difficile* treatments: a systematic review. *Ann Intern Med* 2011; 155: 839–847.
76. Keighley MR, Burdon DW, Arabi Y et al. Randomised controlled trial of vancomycin for pseudomembranous colitis and postoperative diarrhoea. *Br Med J* 1978; 2: 1667–1669.
77. Teasley DG, Gerding DN, Olson MM et al. Prospective randomised trial of metronidazole versus vancomycin for *Clostridium-difficile*-associated diarrhoea and colitis. *Lancet* 1983; 2: 1043–1046.
78. Young GP, Ward PB, Bayley N et al. Antibiotic-associated colitis due to *Clostridium difficile*: double-blind comparison of vancomycin with bacitracin. *Gastroenterology* 1985; 89: 1038.
79. Dudley MN, McLaughlin JC, Carrington G, Frick J, Nightingale CH, Quintiliani R. Oral bacitracin vs vancomycin therapy for *Clostridium difficile*-induced diarrhoea. A randomized double-blind trial. *Arch Intern Med* 1986; 146: 1101–1104.
80. Fekety R, Silva J, Kauffman C, Buggy B, Deery HG. Treatment of antibiotic-associated *Clostridium difficile* colitis with oral vancomycin: comparison of two dosage regimens. *Am J Med* 1989; 86: 15–19.
81. Boero M, Berti E, Morgando A, Verme G. Terapia della colite da *Clostridium difficile*: risultati di uno studio randomizzato aperto rifaximina vs. vancomicina. [Treatment for colitis caused by *Clostridium difficile*: results of a randomized open study of rifaximine vs. vancomycin]. *Microbiol Med* 1990; 5: 74–77.
82. De Lalla F, Nicolini R, Rinaldi E et al. Prospective study of oral teicoplanin versus oral vancomycin for therapy of pseudomembranous colitis and *Clostridium difficile*-associated diarrhoea. *Antimicrob Agents Chemother* 1992; 36: 2192–2196.
83. The Swedish CDAD Study Group. Treatment of *Clostridium difficile* associated diarrhoea and colitis with an oral preparation of teicoplanin; a dose finding study. *Scand J Infect Dis* 1994; 26: 309–316.
84. Wenisch C, Parschalk B, Hasenhüdl M, Hirschl AM, Graninger W. Comparison of vancomycin, teicoplanin, metronidazole, and fusidic acid for the treatment of *Clostridium difficile*-associated diarrhoea. *Clin Infect Dis* 1996; 22: 813–818.
85. Wullt M, Odenholt I. A double-blind randomized controlled trial of fusidic acid and metronidazole for treatment of an initial episode of *Clostridium difficile*-associated diarrhoea. *J Antimicrob Chemother* 2004; 54: 211–216.
86. Musher DM, Logan N, Hamill RJ et al. Nitazoxanide for the treatment of *Clostridium difficile* colitis. *Clin Infect Dis* 2006; 43: 421–427.
87. Lagrotteria D, Holmes S, Smieja M, Smaill F, Lee C. Prospective, randomized inpatient study of oral metronidazole versus oral metronidazole and rifampin for treatment of primary episode of *Clostridium difficile*-associated diarrhoea. *Clin Infect Dis* 2007; 43: 547–552.
88. Zar FA, Bakkanagari SR, Moorthi KMLST, Davis MB. A comparison of vancomycin and metronidazole for the treatment of *Clostridium difficile*-associated diarrhoea, stratified by disease severity. *Clin Infect Dis* 2007; 45: 302–307.
89. Louie T, Miller M, Donskey C, Mullane K, Goldstein EJ. Clinical outcomes, safety, and pharmacokinetics of OPT-80 in a phase 2 trial with patients with *Clostridium difficile* infection. *Antimicrob Agents Chemother* 2008; 53: 223–228.
90. Musher DM, Logan N, Bressler AM, Johnson DP, Rossignol JF. Nitazoxanide versus vancomycin in *Clostridium difficile* infection: a randomized, double-blind study. *Clin Infect Dis* 2009; 48: e41–e46.
91. Cornely OA, Crook DW, Esposito R et al. Fidaxomicin versus vancomycin for infection with *Clostridium difficile* in Europe, Canada, and the USA: a double-blind, non-inferiority, randomised controlled trial. *Lancet Infect Dis* 2012; 12: 281–289.
92. Johnson S, Gerding D, Davidson D et al. Efficacy and safety of oral vancomycin versus oral metronidazole for treatment of *Clostridium difficile*-associated diarrhoea (CDAD): pooled results from two randomized clinical trials. Poster presentation ID 2012. Available at <https://idsa.confex.com/idsa/2012/webprogram/Paper35060.html>
93. Freeman J, Baines SD, Saxton K, Wilcox MH. Effect of metronidazole on growth and toxin production by epidemic *Clostridium difficile* PCR ribotypes 001 and 027 in a human gut model. *J Antimicrob Chemother* 2007; 60: 83–91.
94. Brazier JS, Fawley W, Freeman J, Wilcox MH. Reduced susceptibility of *Clostridium difficile* to metronidazole. *J Antimicrob Chemother* 2001; 48: 741–742.

95. Baines SD, O'Connor R, Freeman J et al. Emergence of reduced susceptibility to metronidazole in *Clostridium difficile*. *J Antimicrob Chemother* 2008; 62: 1046–1052.
96. Moura I, Spigaglia P, Barbanti F, Mastrantonio P. Analysis of metronidazole susceptibility in different *Clostridium difficile* PCR ribotypes. *J Antimicrob Chemother* 2013; 68: 362–365.
97. Johnson S, Sanchez JL, Gerding DN. Metronidazole resistance in *Clostridium difficile*. *Clin Infect Dis* 2000 Aug; 31: 625–626.
98. Pelaez T, Alcalá L, Alonso R, Rodríguez-Creixems M, García-Lechuz JM, Bouza E. Reassessment of *Clostridium difficile* susceptibility to metronidazole and vancomycin. *Antimicrob Agents Chemother* 2002; 46: 1647–1650.
99. Purdell J. Investigation of outcome in cases of *Clostridium difficile* infection due to isolates with reduced susceptibility to metronidazole. In: 21st European Congress of Clinical Microbiology and Infectious Diseases (ECCMID); 2011 in Milan, Italy. Abstract: O499.
100. Bolton RPR, Culshaw MAM. Faecal metronidazole concentrations during oral and intravenous therapy for antibiotic associated colitis due to *Clostridium difficile*. *Gut* 1986; 27: 1169–1172.
101. Al-Nassir WN, Sethi AK, Li Y, Pultz MJ, Riggs MM, Donskey CJ. Both oral metronidazole and oral vancomycin promote persistent overgrowth of vancomycin-resistant enterococci during treatment of *Clostridium difficile*-associated disease. *Antimicrob Agents Chemother* 2008 Jun; 24: 2403–2406.
102. Sethi AK, Nassir Al WN, Nerandzic MM, Donskey CJ. Skin and environmental contamination with vancomycin-resistant enterococci in patients receiving oral metronidazole or oral vancomycin treatment for *Clostridium difficile*-associated disease. *Infect Control Hosp Epidemiol* 2009; 30: 13–17.
103. Miller M, Bernard L, Thompson M et al. Lack of increased colonization with vancomycin-resistant enterococci during preferential use of vancomycin for treatment during an outbreak of healthcare-associated *Clostridium difficile* infection. *Infect Control Hosp Epidemiol* 2010; 31: 710–715.
104. Louie TJ, Cannon K, Byrne B et al. Fidaxomicin preserves the intestinal microbiome during and after treatment of *Clostridium difficile* infection (CDI) and reduces both toxin reexpression and recurrence of CDI. *Clin Infect Dis* 2012; 55 (suppl 2): S132–S142.
105. Nerandzic MM, Mullane K, Miller MA, Babakhani F, Donskey CJ. Reduced acquisition and overgrowth of vancomycin-resistant enterococci and *Candida* species in patients treated with fidaxomicin versus vancomycin for *Clostridium difficile* infection. *Clin Infect Dis* 2012; 55 (suppl 2): S121–S126.
106. De Lalla F, Privitera G, Rinaldi E, Ortisi G, Santoro D, Rizzardini G. Treatment of *Clostridium difficile*-associated disease with teicoplanin. *Antimicrob Agents Chemother* 1989; 33: 1125–1127.
107. Löfmark S, Edlund C, Nord CE. Metronidazole is still the drug of choice for treatment of anaerobic infections. *Clin Infect Dis* 2010; 50 (suppl 1): S16–S23.
108. McGrath NM, Kent-Smith B, Sharp DM. Reversible optic neuropathy due to metronidazole. *Clin Experiment Ophthalmol* 2007; 35: 585–586.
109. Howard-Thompson A, Hurdle AC, Arnold LB, Finch CK, Sands C, Self TH. Intracerebral hemorrhage secondary to a warfarin–metronidazole interaction. *Am J Geriatr Pharmacother* 2008; 6: 33–36.
110. Aradhya S, Manian FA, Hafidh SAS, Bhutto SS, Alpert MA. Significant absorption of oral vancomycin in a patient with *Clostridium difficile* colitis and normal renal function. *South Med J* 2006; 99: 518–520.
111. Weiss K, Allgren RL, Sellers S. Safety analysis of fidaxomicin in comparison with oral vancomycin for *Clostridium difficile* infections. *Clin Infect Dis* 2012; 55 (suppl 2): S110–S115.
112. Iarikov DE, Alexander J, Nambiar S. Hypersensitivity reactions associated with fidaxomicin use. *Clin Infect Dis* 2013, in press.
113. Bartlett JGJ, Tedesco FJF, Shull SS, Lowe BB, Chang TT. Symptomatic relapse after oral vancomycin therapy of antibiotic-associated pseudomembranous colitis. *Gastroenterology* 1980; 78: 431–434.
114. Silva J, Batts DH, Fekety R, Plouffe JF, Rifkin GD, Baird I. Treatment of *Clostridium difficile* colitis and diarrhea with vancomycin. *Am J Med* 1981; 71: 815–822.
115. Cherry RDR, Portnoy DD, Jabbari MM, Daly DSD, Kinnear DGD, Goresky CAC. Metronidazole: an alternate therapy for antibiotic-associated colitis. *Gastroenterology* 1982; 82: 849–851.
116. Bartlett JG. Treatment of antibiotic-associated pseudomembranous colitis. *Clin Infect Dis* 1984; 6 (suppl 1): S235–S241.
117. Olson MM, Shanholtzer CJ, Lee JT, Gerding DN. Ten years of prospective *Clostridium difficile*-associated disease surveillance and treatment at the Minneapolis VA Medical Center, 1982–1991. *Infect Control Hosp Epidemiol* 1994; 15: 371–381.
118. Fernandez A, Anand G, Friedenberg F. Factors associated with failure of metronidazole in *Clostridium difficile*-associated disease. *J Clin Gastroenterol* 2004; 38: 414.
119. Musher DM, Aslam S, Logan N et al. Relatively poor outcome after treatment of *Clostridium difficile* colitis with metronidazole. *Clin Infect Dis* 2005; 40: 1586–1590.
120. Louie TJ. Treating *Clostridium difficile* in the future: what's coming? 45th Interscience Conference on Antimicrobial Agents and Chemotherapy; December 16–19, 2005; Washington, DC. Abstract 1774.
121. Musher DM, Logan N, Mehendiratta V, Melgarejo NA, Garud S, Hamill RJ. *Clostridium difficile* colitis that fails conventional metronidazole therapy: response to nitazoxanide. *J Antimicrob Chemother* 2007; 59: 705–710.
122. Herpers BL, Vlamincx B, Burkhardt O et al. Intravenous tigecycline as adjunctive or alternative therapy for severe refractory *Clostridium difficile* infection. *Clin Infect Dis* 2009; 48: 1732–1735.
123. Rubin DT, Sohi S, Glathar M, Thomas T, Yadron N, Surma BL. Rifaximin is effective for the treatment of *Clostridium difficile* associated diarrhea: results of an open-label pilot study. *Gastroenterol Res Pract* 2011. doi: 10.1155/2011/106978
124. Louie TJ, Gerson M, Grimard D et al. Results of a phase III trial comparing tolevamer, vancomycin and metronidazole in patients with *Clostridium difficile*-associated diarrhea (CDAD). Program and abstracts of the 47th Interscience Conference on Antimicrobial Agents and Chemotherapy, September 17–20, 2007; Chicago, USA. Abstract K-425a.
125. Bouza E, Dryden M, Mohammed R et al. Results of a phase III trial comparing tolevamer, vancomycin and metronidazole in patients with *Clostridium difficile*-associated diarrhoea. Program and abstracts of the 18th European Congress of Clinical Microbiology and Infectious Diseases April 19–22, 2008; Barcelona, Spain. Abstract O464.
126. McFarland LV, Surawicz CM, Greenberg RN et al. A randomized placebo-controlled trial of *Saccharomyces boulardii* in combination with standard antibiotics for *Clostridium difficile* disease. *JAMA* 1994 Jun; 271: 1913–1918.
127. Mogg GA, George RH, Youngs D et al. Randomized controlled trial of colestipol in antibiotic-associated colitis. *Br J Surg* 1982; 69: 137–139.
128. van Dissel JT. Bovine antibody-enriched whey to aid in the prevention of a relapse of *Clostridium difficile*-associated diarrhoea: preclinical and preliminary clinical data. *J Med Microbiol* 2005; 54: 197–205.
129. Numan SC, Veldkamp P, Kuijper EJ, van den Berg RJ, van Dissel JT. *Clostridium difficile*-associated diarrhoea: bovine anti-*Clostridium difficile* whey protein to help aid the prevention of relapses. *Gut* 2007; 56: 888–889.
130. Abougergi MS, Kwon JH. Intravenous immunoglobulin for the treatment of *Clostridium difficile* infection: a review. *Dig Dis Sci* 2011; 56: 19–26.



131. Hempel S, Newberry SJ, Maher AR et al. Probiotics for the prevention and treatment of antibiotic-associated diarrhea: a systematic review and meta-analysis. *JAMA* 2012; 307: 1959–1969.
132. Johnston BC, Ma SSY, Goldenberg JZ et al. Probiotics for the prevention of *Clostridium difficile* associated diarrhea. *Ann Intern Med* 2012; 157: 878–888.
133. Pillai A, Nelson R. Probiotics for treatment of *Clostridium difficile*-associated colitis in adults. *Cochrane Database Syst Rev* 2008; CD004611. doi: 10.1002/14651858.CD004611.pub2
134. Enache-Angoulvant A, Hennequin C. Invasive *Saccharomyces* infection: a comprehensive review. *Clin Infect Dis* 2005; 41: 1559–1568.
135. Muñoz P, Bouza E, Cuenca-Estrella M et al. *Saccharomyces cerevisiae* fungemia: an emerging infectious disease. *Clin Infect Dis* 2005; 40: 1625–1634.
136. Besselink MGH, van Santvoort HC, Buskens E et al. Probiotic prophylaxis in predicted severe acute pancreatitis: a randomised, double-blind, placebo-controlled trial. *Lancet* 2008; 371: 651–659.
137. McFarland LV. Evidence-based review of probiotics for antibiotic-associated diarrhea and *Clostridium difficile* infections. *Anaerobe* 2009; 15: 274–280.
138. Bhangu A, Nepogodiev D, Gupta A, Torrance A, Singh P. West Midlands research collaborative systematic review and meta-analysis of outcomes following emergency surgery for *Clostridium difficile* colitis. *Br J Surg* 2012; 99: 1501–1513.
139. Stewart DB, Hollenbeak CS, Wilson MZ. Is colectomy for fulminant *C. difficile* colitis life saving? A systematic review *Colorectal Dis* 2013; 15: 798–804.
140. Koss K, Clark MA, Sanders DSA, Morton D, Keighley MRB, Goh J. The outcome of surgery in fulminant *Clostridium difficile* colitis. *Colorectal Dis* 2006; 8: 149–154.
141. Chan S, Kelly M, Helme S, Gossage J, Modarai B, Forshaw M. Outcomes following colectomy for *Clostridium difficile* colitis. *Int J Surg* 2009; 7: 78–81.
142. Lee DY, Chung EL, Guend H, Whelan RL, Wedderburn RV, Rose KM. Predictors of mortality after emergency colectomy for *Clostridium difficile* colitis: an analysis of ACS-NSQIP. *Ann Surg* 2013. doi:10.1097/SLA.0b013e31828a8eba.
143. Neal MD, Alverdy JC, Hall DE, Simmons RL, Zuckerbraun BS. Diverting loop ileostomy and colonic lavage: an alternative to total abdominal colectomy for the treatment of severe, complicated *Clostridium difficile* associated disease. *Ann Surg* 2011; 254: 423–437.
144. Cornely OA, Miller MA, Louie TJ, Crook DW, Gorbach SL. Treatment of first recurrence of *Clostridium difficile* infection: fidaxomicin versus vancomycin. *Clin Infect Dis* 2012; 55 (suppl 2): S154–S161.
145. van Nood E, Vrieze A, Nieuwdorp M et al. Duodenal infusion of donor feces for recurrent *Clostridium difficile*. *N Engl J Med* 2013; 368: 407–415.
146. Surawicz CM, Surawicz CM, McFarland LV et al. The search for a better treatment for recurrent *Clostridium difficile* disease: use of high-dose vancomycin combined with *Saccharomyces boulardii*. *Clin Infect Dis* 2000; 31: 1012–1017.
147. Wullt M, Hagslätt M-LJ, Odenholt I. *Lactobacillus plantarum* 299v for the treatment of recurrent *Clostridium difficile*-associated diarrhoea: a double-blind, placebo-controlled trial. *Scand J Infect Dis* 2003; 35: 365–367.
148. Lawrence SJ, Korzenik JR, Mundy LM. Probiotics for recurrent *Clostridium difficile* disease. *J Med Microbiol* 2005; 54: 905–906.
149. Mattila E, Anttila V-J, Broas M et al. A randomized, double-blind study comparing *Clostridium difficile* immune whey and metronidazole for recurrent *Clostridium difficile*-associated diarrhoea: efficacy and safety data of a prematurely interrupted trial. *Scand J Infect Dis* 2008; 40: 702–708.
150. Tedesco FJF, Gordon DD, Fortson WCW. Approach to patients with multiple relapses of antibiotic-associated pseudomembranous colitis. *Am J Gastroenterol* 1985; 80: 867–868.
151. Buggy BP, Fekety R, Silva J Jr. Therapy of relapsing *Clostridium difficile*-associated diarrhea and colitis with the combination of vancomycin and rifampin. *J Clin Gastroenterol* 1987; 9: 155.
152. Johnson S, Schriever C, Galang M, Kelly CP, Gerding DN. Interruption of recurrent *Clostridium difficile*-associated diarrhea episodes by serial therapy with vancomycin and rifaximin. *Clin Infect Dis* 2007; 44: 846–848.
153. Garey KW, Jiang Z-D, Bellard A, DuPont HL. Rifaximin in treatment of recurrent *Clostridium difficile*-associated diarrhea: an uncontrolled pilot study. *J Clin Gastroenterol* 2009; 43: 91–92.
154. Basu PP, Dinani A, Rayapudi K et al. Rifaximin therapy for metronidazole-unresponsive *Clostridium difficile* infection: a prospective pilot trial. *Therap Adv Gastroenterol* 2010; 3: 221–225.
155. Gorbach SL, Chang TW, Goldin B. Successful treatment of relapsing *Clostridium difficile* colitis with *Lactobacillus* GG. *Lancet* 1987; 2: 1519.
156. Biller JA, Katz AJ, Flores AF, Buie TM, Gorbach SL. Treatment of recurrent *Clostridium difficile* colitis with *Lactobacillus* GG. *J Pediatr Gastroenterol Nutr* 1995; 21: 224–226.
157. Bowden TA, Mansberger AR, Lykins LE. Pseudomembranous enterocolitis: mechanism for restoring floral homeostasis. *Am Surg* 1981; 47: 178–183.
158. Tvede M, Rask-Madsen J. Bacteriotherapy for chronic relapsing *Clostridium difficile* diarrhoea in six patients. *Lancet* 1989; 1: 1156.
159. Paterson DLD, Iredell JJ, Whitby MM. Putting back the bugs: bacterial treatment relieves chronic diarrhoea. *Med J Aust* 1994; 160: 232–233.
160. Lund-Tønnesen S, Berstad A, Schreiner A, Midtvedt T. *Clostridium difficile*-associated diarrhea treated with homologous feces. *Tidsskr Nor Laegeforen* 1998; 118: 1027–1030.
161. Faust G, Langelier D, Haddad H, Menard DB. Treatment of recurrent pseudomembranous colitis (RPMC) with stool transplantation (ST): report of six (6) cases. 41st annual meeting of the Canadian Association of Gastroenterology in conjunction with the Canadian Association for the Study of the Liver, 2002; Montreal, Quebec, Canada: Abstract 002.
162. Aas J, Gessert CE, Bakken JS. Recurrent *Clostridium difficile* colitis: case series involving 18 patients treated with donor stool administered via a nasogastric tube. *Clin Infect Dis* 2003; 36: 580–585.
163. Jorup-Rönström C, Håkanson A, Persson AK, Midtvedt T, Norin E. Feces culture successful therapy in *Clostridium difficile* diarrhea. *Lakartidningen* 2006; 103: 3603–3605.
164. Brandt LJ, Reddy SS. Fecal microbiota transplantation for recurrent *Clostridium difficile* infection. *J Clin Gastroenterol* 2011; 45 (suppl): S159–S167.
165. Wettstein A, Borody TJ, Leis S. Fecal bacteriotherapy: an effective treatment for relapsing symptomatic *Clostridium difficile* infection. 15th United European Gastroenterology Week; 2007, October 27–31, Paris, France: Abstract G-67]
166. Nieuwdorp M, van Nood E, Speelman P et al. Behandeling van recidiverende *Clostridium difficile*-geassocieerde diarree met een suspensie van donorfeces. *Ned Tijdschr Geneesk* 2008; 152: 1927–1932.
167. Rubin TA, Gessert CE, Aas J. Stool transplantation for older patients with *Clostridium difficile* infection. *J Am Geriatr Soc* 2009; 57: 2386.
168. MacConnachie AA, Fox R, Kennedy DR, Seaton RA. Faecal transplant for recurrent *Clostridium difficile*-associated diarrhoea: a UK case series. *QJM* 2009; 102: 781–784.
169. Arkkila PE, Uusitalo-Seppälä R, Lehtola L, Moilanen V, Ristikankare M, Mattila EJ. Fecal bacteriotherapy for recurrent *Clostridium difficile* infection. *Gastroenterol* 2010; 138: S5.
170. Rohlke F, Surawicz CM, Stollman N. Fecal flora reconstitution for recurrent *Clostridium difficile* infection: results and methodology. *J Clin Gastroenterol* 2010; 44: 567–570.

171. Silverman MS, Davis I, Pillai DR. Success of self-administered home fecal transplantation for chronic *Clostridium difficile* infection. *Clin Gastroenterol Hepatol* 2010; 8: 471–473.
172. Mellow MHM, Kanatzar AA. Colonoscopic fecal bacteriotherapy in the treatment of recurrent *Clostridium difficile* infection – results and follow-up. *J Okla State Med Assoc* 2011; 104: 89–91.
173. Yoon SS, Brandt LJ. Treatment of refractory/recurrent *C. difficile*-associated disease by donated stool transplanted via colonoscopy: a case series of 12 patients. *J Clin Gastroenterol* 2010; 44: 562–566.
174. Garborg K, Waagsbo B, Stallemo A, Matre J, Sundy A. Results of faecal donor instillation therapy for recurrent *Clostridium difficile*-associated diarrhoea. *Scand J Infect Dis* 2010; 42: 857–861.
175. Kelly CR, de Leon L, Jasutkar N. Fecal microbiota transplantation for relapsing *Clostridium difficile* infection in 26 patients: methodology and results. *J Clin Gastroenterol* 2012; 46: 145.
176. Brandt RJ, Aroniadis OC, Mellow M, et al. Long-term follow-up of colonoscopic fecal microbiota transplant for recurrent *Clostridium difficile* infection. *Am J Gastroenterol* 2012; 107: 1079–1087.
177. Kassam Z, Hundal R, Marshall JK, Lee CH. Fecal transplant via retention enema for refractory or recurrent *Clostridium difficile* infection. *Arch Intern Med* 2012; 172: 191–193.
178. Hamilton MJ, Olson MM, Weingarden AR et al. Standardized frozen preparation for transplantation of fecal microbiota for recurrent *Clostridium difficile* infection. *Am J Gastroenterol* 2012; 107: 761–767.
179. Mattila E, Seppälä RU, Wuorela M et al. Fecal transplantation, through colonoscopy, is effective therapy for recurrent *Clostridium difficile* infection. *Gastroenterology* 2012; 142: 490–496.
180. Leung DY, Kelly CP, Boguniewicz M, Pothoulakis C, LaMont JT, Flores A. Treatment with intravenously administered gamma globulin of chronic relapsing colitis induced by *Clostridium difficile* toxin. *J Pediatr* 1991; 118: 633–637.
181. Beales ILP. Intravenous immunoglobulin for recurrent *Clostridium difficile* diarrhoea. *Gut* 2002; 51: 456.
182. McPherson S, Rees CJ, Ellis R, Soo S, Panter SJ. Intravenous immunoglobulin for the treatment of severe, refractory, and recurrent *Clostridium difficile* diarrhea. *Dis Colon Rectum* 2006; 49: 640–645.
183. Juang P, Skledar SJ, Zgheib NK et al. Clinical outcomes of intravenous immune globulin in severe *Clostridium difficile*-associated diarrhea. *Am J Infect Control* 2007; 35: 131–137.
184. Abougergi MS, Broor A, Cui W, Jaar BG. Intravenous immunoglobulin for the treatment of severe *Clostridium difficile* colitis: an observational study and review of the literature. *J Hosp Med* 2010; 5: E1–E9.
185. Bakken JS. Fecal bacteriotherapy for recurrent *Clostridium difficile* infection. *Anaerobe* 2009; 15: 285–289.
186. Landy J, Al-Hassi HO, McLaughlin SD et al. Review article: faecal transplantation therapy for gastrointestinal disease. *Aliment Pharmacol Ther* 2011; 34: 409–415.
187. Kassam Z, Lee CH, Yuan Y, Hunt RH. Fecal microbiota transplantation for *Clostridium difficile* infection: systematic review and meta-analysis. *Am J Gastroenterol* 2013; 108: 500–508.
188. Rohlke F, Stollman N. Fecal microbiota transplantation in relapsing *Clostridium difficile* infection. *Therap Adv Gastroenterol* 2012; 5: 403–420.
189. Guo BB, Harstall CC, Louie TT, van Zanten SSV, Dieleman LAL. Systematic review: faecal transplantation for the treatment of *Clostridium difficile*-associated disease. *Aliment Pharmacol Ther* 2012; 35: 865–875.
190. Gough E, Shaikh H, Manges AR. Systematic review of intestinal microbiota transplantation (fecal bacteriotherapy) for recurrent *Clostridium difficile* infection. *Clin Infect Dis* 2011; 53: 994–1002.
191. Van Nood E, Speelman P, Kuijper EJ, Keller JJ. Struggling with recurrent *Clostridium difficile* infections: is donor faeces the solution? *Euro Surveill* 2009; 14: pii.
192. Friedenberg F, Fernandez A, Kaul V, Niami P, Levine GM. Intravenous metronidazole for the treatment of *Clostridium difficile* colitis. *Dis Colon Rectum* 2001; 44: 1176–1180.
193. McFarland LV. Alternative treatments for *Clostridium difficile* disease: what really works? *J Med Microbiol* 2005; 54: 101–111.
194. Musgrave CR, Bookstaver PB, Sutton SS, Miller AD. Use of alternative or adjuvant pharmacologic treatment strategies in the prevention and treatment of *Clostridium difficile* infection. *Int J Infect Dis* 2011; 15: e438–e448.
195. Larson KC, Belliveau PP, Spooner LM. Tigecycline for the treatment of severe *Clostridium difficile* infection. *Ann Pharmacother* 2011; 45: 1005–1010.