

FIDAXOMICIN- fidaxomicin tablet, film coated
Teva Pharmaceuticals, Inc.

HIGHLIGHTS OF PRESCRIBING INFORMATION

These highlights do not include all the information needed to use FIDAXOMICIN TABLETS safely and effectively. See full prescribing information for FIDAXOMICIN TABLETS.

FIDAXOMICIN tablets, for oral use
Initial U.S. Approval: 2011

INDICATIONS AND USAGE

Fidaxomicin tablets are a macrolide antibacterial indicated in adult patients for the treatment of *C. difficile*-associated diarrhea. (1.1)

To reduce the development of drug-resistant bacteria and maintain the effectiveness of fidaxomicin and other antibacterial drugs, fidaxomicin should be used only to treat infections that are proven or strongly suspected to be caused by *C. difficile*. (1.2)

DOSAGE AND ADMINISTRATION

- Fidaxomicin tablets are administered orally with or without food. (2.1)
- Adults
 - One 200 mg tablet orally twice daily for 10 days. (2.2)

DOSAGE FORMS AND STRENGTHS

- Film-coated tablets: 200 mg (3)

CONTRAINDICATIONS

Fidaxomicin is contraindicated in patients who have known hypersensitivity to fidaxomicin or any other ingredient in fidaxomicin tablets. (4)

WARNINGS AND PRECAUTIONS

- Acute hypersensitivity reactions (angioedema, dyspnea, pruritus, and rash) have been reported. If a severe hypersensitivity reaction occurs, discontinue fidaxomicin. (5.1)
- Fidaxomicin is not expected to be effective for the treatment of other types of infections due to minimal systemic absorption of fidaxomicin.
Fidaxomicin should only be used for the treatment of *C. difficile*-associated diarrhea. (5.2)
- Development of drug-resistant bacteria: Only use fidaxomicin for infection proven or strongly suspected to be caused by *C. difficile*. (5.3)

ADVERSE REACTIONS

The most common adverse reactions in adults (incidence $\geq 2\%$) are nausea, vomiting, abdominal pain, gastrointestinal hemorrhage, anemia, and neutropenia. (6)

To report SUSPECTED ADVERSE REACTIONS, contact Teva at 1-888-838-2872 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

USE IN SPECIFIC POPULATIONS

Pediatrics: The safety and effectiveness of fidaxomicin have not been established in pediatric patients younger than 6 months of age. (8.4)

Pediatric use information is approved for Cubist Pharmaceuticals LLC's DIFICID® (fidaxomicin) tablets. However, due to Cubist Pharmaceuticals LLC's marketing exclusivity rights, this drug product is not labeled with that information.

See 17 for PATIENT COUNSELING INFORMATION and FDA-approved patient labeling.

Revised: 5/2024

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FULL PRESCRIBING INFORMATION

1 INDICATIONS AND USAGE

1.1 *Clostridioides difficile*-Associated Diarrhea

Fidaxomicin tablets are indicated in adult patients for the treatment of *C. difficile*-associated diarrhea (CDAD).

1.2 Usage

To reduce the development of drug-resistant bacteria and maintain the effectiveness of fidaxomicin tablets and other antibacterial drugs, fidaxomicin tablets should be used only to treat infections that are proven or strongly suspected to be caused by *C. difficile*. When culture and susceptibility information are available, they should be considered in selecting or modifying antibacterial therapy. In the absence of such data, local epidemiology and susceptibility patterns may contribute to the empiric selection of therapy.

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2 DOSAGE AND ADMINISTRATION

2.1 Important Administration Instructions

Fidaxomicin tablets are available for oral administration as 200 mg tablets. Fidaxomicin tablets are administered orally with or without food.

2.2 Adult Patients

The recommended dosage for adults is one 200 mg fidaxomicin tablet orally twice daily for 10 days.

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3 DOSAGE FORMS AND STRENGTHS

Film-coated tablets: Each white, film-coated, modified capsule shape tablet debossed with A205 on one side and plain on the other side contains 200 mg of fidaxomicin.

4 CONTRAINDICATIONS

Fidaxomicin is contraindicated in patients who have known hypersensitivity to fidaxomicin or any other ingredient in fidaxomicin tablets [see *Warnings and Precautions (5.1)*].

5 WARNINGS AND PRECAUTIONS

5.1 Hypersensitivity Reactions

Acute hypersensitivity reactions, including dyspnea, rash, pruritus, and angioedema of the mouth, throat, and face have been reported with fidaxomicin. If a severe hypersensitivity reaction occurs, fidaxomicin should be discontinued and appropriate therapy should be instituted.

Some patients with hypersensitivity reactions to fidaxomicin also reported a history of allergy to other macrolides. Physicians prescribing fidaxomicin to patients with a known macrolide allergy should be aware of the possibility of hypersensitivity reactions.

5.2 Not for Use in Infections Other than *C. difficile*-Associated Diarrhea

Fidaxomicin is not expected to be effective for the treatment of other types of infections due to minimal systemic absorption of fidaxomicin [see *Clinical Pharmacology (12.3)*]. Fidaxomicin has not been studied for the treatment of infections other than CDAD. Fidaxomicin should only be used for the treatment of CDAD.

5.3 Development of Drug-Resistant Bacteria

Prescribing fidaxomicin in the absence of proven or strongly suspected *C. difficile* infection is unlikely to provide benefit to the patient and increases the risk of the development of drug-resistant bacteria.

6 ADVERSE REACTIONS

6.1 Clinical Trials Experience

Because clinical trials are conducted under widely varying conditions, adverse reaction rates observed in the clinical trials of a drug cannot be directly compared to rates in the clinical trials of another drug and may not reflect the rates observed in practice.

Adults

The safety of fidaxomicin 200 mg tablets taken twice a day for 10 days was evaluated in 564 adult patients with CDAD in two active-controlled trials with 86.7% of patients receiving a full course of treatment.

Thirty-three adult patients receiving fidaxomicin (5.9%) withdrew from trials as a result of adverse reactions (AR). The types of AR resulting in withdrawal from the study varied considerably. Vomiting was the primary adverse reaction leading to discontinuation of dosing; this occurred at an incidence of 0.5% in both the fidaxomicin and vancomycin patients in Phase 3 trials. The most common selected adverse reactions occurring in $\geq 2\%$ of adult patients treated with fidaxomicin are listed in Table 2.

Table 2: Selected Adverse Reactions with an Incidence of $\geq 2\%$ Reported in Fidaxomicin-Treated Adult Patients in Controlled Trials

System Organ Class Adverse Reaction	Fidaxomicin (N=564)	Vancomycin (N=583)
	n (%)	n (%)
Blood and Lymphatic System Disorders		
Anemia	14 (2%)	12 (2%)
Neutropenia	14 (2%)	6 (1%)
Gastrointestinal Disorders		
Nausea	62 (11%)	66 (11%)
Vomiting	41 (7%)	37 (6%)
Abdominal Pain	33 (6%)	23 (4%)
Gastrointestinal Hemorrhage	20 (4%)	12 (2%)

The following adverse reactions were reported in $<2\%$ of adult patients taking fidaxomicin tablets in controlled trials:

Gastrointestinal Disorders: abdominal distension, abdominal tenderness, dyspepsia, dysphagia, flatulence, intestinal obstruction, megacolon

Investigations: increased blood alkaline phosphatase, decreased blood bicarbonate, increased hepatic enzymes, decreased platelet count

Metabolism and Nutrition Disorders: hyperglycemia, metabolic acidosis

Skin and Subcutaneous Tissue Disorders: drug eruption, pruritus, rash

6.2 Postmarketing Experience

The following adverse reactions have been identified during post-approval use of fidaxomicin. Because these reactions are reported voluntarily from a population of uncertain size, it is not always possible to reliably estimate their frequency or establish a causal relationship to drug exposure.

Hypersensitivity reactions (dyspnea, angioedema, rash, pruritus)

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7 DRUG INTERACTIONS

Fidaxomicin and its main metabolite, OP-1118, are substrates of the efflux transporter, P-glycoprotein (P-gp), which is expressed in the gastrointestinal tract.

7.1 Cyclosporine

Cyclosporine is an inhibitor of multiple transporters, including P-gp. When cyclosporine was coadministered with fidaxomicin, plasma concentrations of fidaxomicin and OP-1118 were significantly increased but remained in the ng/mL range [see *Clinical Pharmacology (12.3)*]. Concentrations of fidaxomicin and OP-1118 may also be decreased at the site of action (i.e., gastrointestinal tract) via P-gp inhibition; however, concomitant P-gp inhibitor use had no attributable effect on safety or treatment outcome of fidaxomicin-treated adult patients in controlled clinical trials. Based on these results, fidaxomicin may be coadministered with P-gp inhibitors and no dose adjustment is recommended.

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Risk Summary

The limited available data on use of fidaxomicin in pregnant women are insufficient to inform any drug-associated risk for major birth defects, miscarriage or adverse maternal or fetal outcomes. Embryo-fetal reproduction studies in rats and rabbits dosed intravenously during organogenesis revealed no evidence of harm to the fetus at fidaxomicin and OP-1118 (its main metabolite) exposures 65-fold or higher than the clinical exposure at the fidaxomicin recommended dose [see *Data*].

The estimated background risk of major birth defects and miscarriage for the indicated populations is unknown. All pregnancies have a background risk of birth defect, loss, or other adverse outcomes. In the U.S. general population, the estimated background risk of major birth defects and miscarriage in clinically recognized pregnancies is 2% to 4% and 15% to 20%, respectively.

Data

Animal Data

In pregnant rats, fidaxomicin was administered intravenously at doses of 4, 8, and 15 mg/kg/day from gestation day 6 through 17 (during the period of organogenesis). No embryo/fetal effects were noted in this study at exposures (AUC) 193-fold higher for fidaxomicin, and 65-fold higher for OP-1118 than the clinical exposure at the fidaxomicin recommended dose.

In pregnant rabbits, fidaxomicin was administered intravenously at doses of 2, 4, and 7.5 mg/kg/day from gestation day 6 through 18 (during the period of organogenesis). No embryo/fetal effects were noted in this study at exposures 66-fold higher for fidaxomicin, and 245-fold higher for OP-1118 than the clinical exposure at the fidaxomicin recommended dose.

8.2 Lactation

Risk Summary

There is no information on the presence of fidaxomicin or its main metabolite, OP-1118, in human milk, the effects on the breastfed infant, or the effects on milk production. The developmental and health benefits of breastfeeding should be considered along with the mother's clinical need for fidaxomicin and any potential adverse effects on the breastfed infant from fidaxomicin or from the underlying maternal condition.

8.4 Pediatric Use

The safety and effectiveness of fidaxomicin have not been established in pediatric patients younger than 6 months of age.

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8.5 Geriatric Use

Of the total number of patients in controlled trials of fidaxomicin, 50% were 65 years of age and over, while 31% were 75 and over. No overall differences in safety or effectiveness of fidaxomicin compared to vancomycin were observed between these subjects and younger subjects.

In controlled trials, elderly patients (≥65 years of age) had higher plasma concentrations of fidaxomicin and its main metabolite, OP-1118, versus non-elderly patients (<65 years of age) [see *Clinical Pharmacology* (12.3)]. However, greater exposures in elderly patients were not considered to be clinically significant. No dose adjustment is recommended for elderly patients.

10 OVERDOSAGE

No cases of acute overdose have been reported in humans. No drug-related adverse effects were seen in dogs dosed with fidaxomicin tablets at 9,600 mg/day (over 100 times the human dose, scaled by weight) for 3 months.

11 DESCRIPTION

Fidaxomicin is a macrolide antibacterial drug for oral administration. Its CAS chemical name is Oxacyclooctadeca-3,5,9,13,15-pentaen-2-one, 3-[[[6-deoxy-4-O-(3,5-dichloro-2-ethyl-4,6-dihydroxybenzoyl)-2-O-methyl-β-D-mannopyranosyl]oxy]methyl]-12-[[6-deoxy-5-C-methyl-4-O-(2-methyl-1-oxopropyl)-β-D-lyxo-hexopyranosyl]oxy]-11-ethyl-8-hydroxy-18-[(1R)-1-hydroxyethyl]-9,13,15-trimethyl-, (3E,5E,8S,9E,11S,12R,13E,15E,18S)-. The molecular formula is C₅₂H₇₄Cl₂O₁₈ and the molecular weight is 1058.04 g/mol. The structural formula of fidaxomicin is shown in Figure 1.

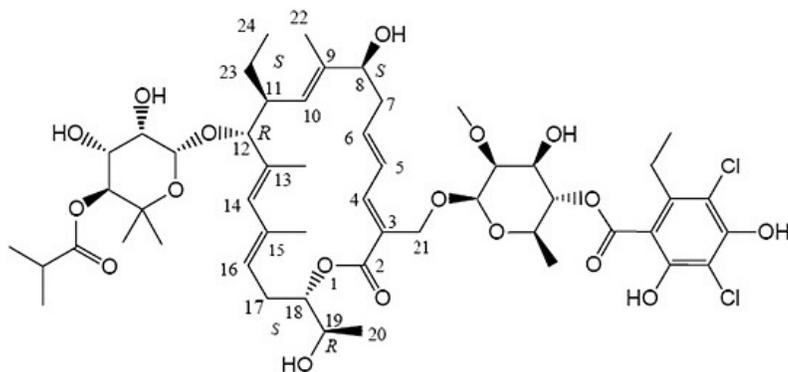


Figure 1: Structural Formula of Fidaxomicin

Fidaxomicin tablets are film-coated and contain 200 mg of fidaxomicin per tablet and the following inactive ingredients: butylated hydroxytoluene, hydroxypropyl cellulose, lecithin (soy), magnesium stearate, microcrystalline cellulose, polyethylene glycol 3350, polyvinyl alcohol-part hydrolyzed, pregelatinized corn starch, sodium starch glycolate Type A, talc, and titanium dioxide.

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

Fidaxomicin is an antibacterial drug [see *Microbiology (12.4)*].

12.2 Pharmacodynamics

Fidaxomicin acts locally in the gastrointestinal tract on *C. difficile*. In a dose-ranging trial (N=48) of fidaxomicin using 50 mg, 100 mg, and 200 mg twice daily for 10 days, a dose-response relationship was observed for efficacy.

12.3 Pharmacokinetics

The pharmacokinetic parameters of fidaxomicin and its main metabolite OP-1118 following a single dose of 200 mg in healthy adult males (N=14) are summarized in Table 4.

Table 4: Mean (± Standard Deviation) Pharmacokinetic Parameters of Fidaxomicin 200 mg in Healthy Adult Males

Parameter	Fidaxomicin		OP-1118	
	N	Value	N	Value
C _{max} (ng/mL)	14	5.20 ± 2.81	14	12.0 ± 6.06
T _{max} (h)*	14	2.00 (1.00 to 5.00)	14	1.02 (1.00 to 5.00)
AUC _{0-t} (ng-h/mL)	14	48.3 ± 18.4	14	103 ± 39.4
AUC _{0-∞} (ng-h/mL)	9	62.9 ± 19.5	10	118 ± 43.3
t _{1/2} (h)	9	11.7 ± 4.80	10	11.2 ± 3.01

* T_{max}, reported as median (range).

C_{max}, maximum observed concentration; T_{max}, time to maximum observed concentration; AUC_{0-t}, area under the concentration-time curve from time 0 to the last measured concentration; AUC_{0-∞}, area under the concentration-time curve from time 0 to infinity; t_{1/2}, elimination half-life

Absorption

Fidaxomicin has minimal systemic absorption following oral administration, with plasma concentrations of fidaxomicin and OP-1118 in the ng/mL range at the therapeutic dose. In fidaxomicin-treated patients from controlled trials, plasma concentrations of fidaxomicin and OP-1118 obtained within the T_{max} window (1 to 5 hours) were approximately 2- to 6-fold higher than C_{max} values in healthy adults. Following administration of fidaxomicin tablets 200 mg twice daily for 10 days, OP-1118 plasma concentrations within the T_{max} window were approximately 50% to 80% higher than on Day 1, while concentrations of fidaxomicin were similar on Days 1 and 10.

In a food-effect study involving administration of fidaxomicin to healthy adults (N=28) with a high-fat meal versus under fasting conditions, C_{max} of fidaxomicin and OP-1118 decreased by 21.5% and 33.4%, respectively, while AUC_{0-t} remained unchanged. This decrease in C_{max} is not considered clinically significant, and thus, fidaxomicin may be administered with or without food.

Distribution

Fidaxomicin is mainly confined to the gastrointestinal tract following oral administration. In selected patients (N=8) treated with fidaxomicin 200 mg twice daily for 10 days from controlled trials, fecal concentrations of fidaxomicin and OP-1118 obtained within 24 hours of the last dose ranged from 639 to 2710 mcg/g and 213 to 1210 mcg/g, respectively. In contrast, plasma concentrations of fidaxomicin and OP-1118 within the T_{max} window (1 to 5 hours) ranged 2 to 179 ng/mL and 10 to 829 ng/mL, respectively.

Elimination

Metabolism

Fidaxomicin is primarily transformed by hydrolysis at the isobutyryl ester to form its main and microbiologically active metabolite, OP-1118. Metabolism of fidaxomicin and formation of OP-1118 are not dependent on cytochrome P450 (CYP) enzymes.

At the therapeutic dose, OP-1118 was the predominant circulating compound in healthy adults, followed by fidaxomicin.

Excretion

Fidaxomicin is mainly excreted in feces. In one trial of healthy adults (N=11), more than 92% of the dose was recovered in the stool as fidaxomicin and OP-1118 following single doses of 200 mg and 300 mg. In another trial of healthy adults (N=6), 0.59% of the dose was recovered in urine as OP-1118 only following a single dose of 200 mg.

Specific Populations

Geriatric Patients

In controlled trials of patients treated with fidaxomicin 200 mg twice daily for 10 days, mean and median values of fidaxomicin and OP-1118 plasma concentrations within the T_{max} window (1 to 5 hours) were approximately 2- to 4-fold higher in elderly patients (≥ 65 years of age) versus non-elderly patients (< 65 years of age). Despite greater exposures in elderly patients, fidaxomicin and OP-1118 plasma concentrations remained in the ng/mL range [see *Use in Specific Populations (8.5)*].

Pediatric Patients

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Male and Female Patients

Plasma concentrations of fidaxomicin and OP-1118 within the T_{max} window (1 to 5 hours) did not vary by gender in patients treated with fidaxomicin 200 mg twice daily for 10 days from controlled trials. No dose adjustment is recommended based on gender.

Patients with Renal Impairment

In controlled trials of patients treated with fidaxomicin 200 mg twice daily for 10 days, plasma concentrations of fidaxomicin and OP-1118 within the T_{max} window (1 to 5 hours) did not vary by severity of renal impairment (based on creatinine clearance) between mild (51 to 79 mL/min), moderate (31 to 50 mL/min), and severe (≤ 30 mL/min) categories. No dose adjustment is recommended based on renal function.

Patients with Hepatic Impairment

The impact of hepatic impairment on the pharmacokinetics of fidaxomicin has not been evaluated. Because fidaxomicin and OP-1118 do not appear to undergo significant hepatic metabolism, elimination of fidaxomicin and OP-1118 is not expected to be significantly affected by hepatic impairment.

Drug Interaction Studies

In vivo studies were conducted to evaluate intestinal drug-drug interactions of fidaxomicin as a P-gp substrate, P-gp inhibitor, and inhibitor of major CYP enzymes expressed in the gastrointestinal tract (CYP3A4, CYP2C9, and CYP2C19).

Table 5 summarizes the impact of a coadministered drug (P-gp inhibitor) on the pharmacokinetics of fidaxomicin [see *Drug Interactions (7.1)*].

Table 5: Pharmacokinetic Parameters of Fidaxomicin and OP-1118 in the Presence of a Coadministered Drug

Parameter	Cyclosporine 200 mg + Fidaxomicin 200 mg* (N=14)		Fidaxomicin 200 mg Alone (N=14)		Mean Ratio of Parameters With/Without Coadministered Drug (90% CI [†]) No Effect = 1.00
	N	Mean	N	Mean	
Fidaxomicin					
C_{max} (ng/mL)	14	19.4	14	4.67	4.15 (3.23 to 5.32)
$AUC_{0-\infty}$ (ng- h/mL)	8	114	9	59.5	1.92 (1.39 to 2.64)
OP-1118					
C_{max} (ng/mL)	14	100	14	10.6	9.51 (6.93 to 13.05)
$AUC_{0-\infty}$ (ng- h/mL)	12	438	10	106	4.11 (3.06 to 5.53)

* Cyclosporine was administered 1 hour before fidaxomicin.

† CI - confidence interval

Fidaxomicin had no significant impact on the pharmacokinetics of the following coadministered drugs: digoxin (P-gp substrate), midazolam (CYP3A4 substrate), warfarin (CYP2C9 substrate), and omeprazole (CYP2C19 substrate). No dose adjustment is warranted when fidaxomicin is coadministered with substrates of P-gp or CYP enzymes.

12.4 Microbiology

Mechanism of Action

Fidaxomicin is a fermentation product obtained from the Actinomycete *Dactylosporangium aurantiacum*. Fidaxomicin is a macrolide antibacterial drug that inhibits RNA synthesis by binding to RNA polymerases. Fidaxomicin is bactericidal against *C. difficile* *in vitro*, and demonstrates a post-antibiotic effect vs. *C. difficile* of 6 to 10 hrs.

Resistance

Fidaxomicin demonstrates no *in vitro* cross-resistance with other classes of antibacterial drugs. *In vitro* studies indicate a low frequency of spontaneous resistance to fidaxomicin in *C. difficile* (ranging from $<1.4 \times 10^{-9}$ to 12.8×10^{-9}). A specific mutation (Val-1143-Gly) in the beta subunit of RNA polymerase is associated with reduced susceptibility to fidaxomicin. This mutation was created in the laboratory and seen during clinical trials in a *C. difficile* isolate obtained from an adult subject treated with fidaxomicin who had recurrence of CDAD. The fidaxomicin minimum inhibitory concentration (MIC) of the *C. difficile* isolate from this subject increased from a baseline of 0.06 mcg/mL to 16 mcg/mL at the time of CDAD recurrence.

Interaction with Other Antimicrobials

Fidaxomicin and its main metabolite OP-1118 do not exhibit any antagonistic interaction with other classes of antibacterial drugs. Synergistic interactions of fidaxomicin and OP-1118 have been observed *in vitro* with rifampin and rifaximin against *C. difficile*.

Antimicrobial Activity

Fidaxomicin has been shown to be active against most isolates of *Clostridioides* (formerly *Clostridium*) *difficile*, both *in vitro* and in clinical infections [see *Indications and Usage (1)*].

Susceptibility Testing

For specific information regarding susceptibility test interpretive criteria, and associated test methods and quality control standards recognized by FDA for this drug, please see: <https://www.fda.gov/STIC>.

13 NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

Long-term carcinogenicity studies have not been conducted to evaluate the carcinogenic potential of fidaxomicin.

Neither fidaxomicin nor OP-1118 was mutagenic in the Ames assay. Fidaxomicin was also negative in the rat micronucleus assay. However, fidaxomicin was clastogenic in Chinese hamster ovary cells.

Fidaxomicin did not affect the fertility of male and female rats at intravenous doses of 6.3 mg/kg. The exposure (AUC_{0-t}) was approximately 100 times that in humans.

14 CLINICAL STUDIES

14.1 Clinical Studies of Fidaxomicin in Adult Patients with CDAD

In two randomized, double-blinded trials, a non-inferiority design was utilized to demonstrate the efficacy of fidaxomicin (200 mg tablets twice daily for 10 days) compared to vancomycin (125 mg four times daily for 10 days) in adults with CDAD.

Enrolled patients were 18 years of age or older and received no more than 24 hours of pretreatment with vancomycin or metronidazole. CDAD was defined by >3 unformed bowel movements (or >200 mL of unformed stool for subjects having rectal collection devices) in the 24 hours before randomization, and presence of either *C. difficile* toxin A or B in the stool within 48 hours of randomization. Enrolled patients had either no prior CDAD history or only one prior CDAD episode in the past three months. Subjects with

life-threatening/fulminant infection, hypotension, septic shock, peritoneal signs, significant dehydration, or toxic megacolon were excluded.

The demographic profile and baseline CDAD characteristics of enrolled subjects were similar in the two trials. Patients had a median age of 64 years, were mainly white (90%), female (58%), and inpatients (63%). The median number of bowel movements per day was 6, and 37% of subjects had severe CDAD (defined as 10 or more unformed bowel movements per day or WBC \geq 15000/mm³). Diarrhea alone was reported in 45% of patients and 84% of subjects had no prior CDAD episode.

The primary efficacy endpoint was the clinical response rate at the end of treatment, based upon improvement in diarrhea or other symptoms such that, in the investigator's judgment, further CDAD treatment was not needed. An additional efficacy endpoint was sustained clinical response 25 days after the end of treatment. Sustained response was evaluated only for patients who were clinical successes at the end of treatment. Sustained response was defined as clinical response at the end of treatment, and survival without proven or suspected CDAD recurrence through 25 days beyond the end of treatment.

The results for clinical response at the end of treatment in both trials, shown in Table 6, indicate that fidaxomicin is non-inferior to vancomycin based on the 95% confidence interval (CI) lower limit being greater than the non-inferiority margin of -10%.

The results for sustained clinical response at the end of the follow-up period, also shown in Table 6, indicate that fidaxomicin is superior to vancomycin on this endpoint. Since clinical success at the end of treatment and mortality rates were similar across treatment arms (approximately 6% in each group), differences in sustained clinical response were due to lower rates of proven or suspected CDAD during the follow-up period in fidaxomicin patients.

Table 6: Clinical Response Rates at End-of-Treatment and Sustained Response at 25 days Post-Treatment in Adult Patients

	Clinical Response at End of Treatment			Sustained Response at 25 days Post-Treatment		
	Fidaxomicin % (N)	Vancomycin % (N)	Difference (95% CI)*	Fidaxomicin % (N)	Vancomycin % (N)	Difference (95% CI)*
Trial 1	88% (N=289)	86% (N=307)	2.6% (-2.9%, 8.0%)	70% (N=289)	57% (N=307)	12.7% (4.4%, 20.9%)
Trial 2	88% (N=253)	87% (N=256)	1.0% (-4.8%, 6.8%)	72% (N=253)	57% (N=256)	14.6% (5.8%, 23.3%)

* Confidence interval (CI) was derived using Wilson's score method. Approximately 5%-9% of the data in each trial and treatment arm were missing sustained response information and were imputed using multiple imputation method.

Restriction Endonuclease Analysis (REA) was used to identify *C. difficile* baseline isolates in the BI group, isolates associated with increasing rates and severity of CDAD in the US in the years prior to the clinical trials. Similar rates of clinical response at the end of treatment and proven or suspected CDAD during the follow-up period were seen in fidaxomicin-treated and vancomycin-treated patients infected with a BI isolate. However, fidaxomicin did not demonstrate superiority in sustained clinical response when compared with vancomycin (Table 7).

Table 7: Sustained Clinical Response at 25 Days after Treatment by *C. difficile* REA Group at Baseline in Adult Patients

Trial 1			
Initial <i>C. difficile</i> Group	Fidaxomicin n/N (%)	Vancomycin n/N (%)	Difference (95% CI)*
BI Isolates	44/76 (58%)	52/82 (63%)	-5.5% (-20.3%, 9.5%)
Non-BI Isolates	105/126 (83%)	87/131 (66%)	16.9% (6.3%, 27.0%)
Trial 2			
Initial <i>C. difficile</i> Group	Fidaxomicin n/N (%)	Vancomycin n/N (%)	Difference (95% CI)*

BI Isolates	42/65 (65%)	31/60 (52%)	12.9% (-4.2%, 29.2%)
Non-BI Isolates	109/131 (83%)	77/121 (64%)	19.6% (8.7%, 30.0%)

* Interaction test between the effect on sustained response rate and BI versus non-BI isolates using logistic regression (p-values: trial 1: 0.009; trial 2: 0.29). Approximately 25% of the mITT population were missing data for REA group. Confidence intervals (CI) were derived using Wilson's score method.

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16 HOW SUPPLIED/STORAGE AND HANDLING

16.1 How Supplied

Fidaxomicin tablets are supplied as follows:

200 mg - Each white, film-coated, modified capsule shape tablet, debossed with A205 on one side and plain on the other side contains 200 mg of fidaxomicin. Tablets are supplied as bottles of 20 tablets NDC 0480-2596-34.

Store in the original bottle.

16.2 Storage

Store at 20° to 25°C (68° to 77°F); excursions permitted to 15° to 30°C (59° to 86°F) [see USP Controlled Room Temperature].

17 PATIENT COUNSELING INFORMATION

Advise the patient to read the FDA-approved patient labeling (Patient Information).

Administration with Food

Inform patients and caregivers that fidaxomicin tablets may be taken with or without food.

Antibacterial Resistance

Patients should be counseled that antibacterial drugs, including fidaxomicin tablets, should only be used to treat bacterial infections. They do not treat viral infections (e.g., the common cold). When fidaxomicin tablets are prescribed to treat a *C. difficile* infection, patients should be told that, although it is common to feel better early in the course of therapy, the medication should be taken exactly as directed. Skipping doses or not completing the full course of therapy may (1) decrease the effectiveness of the immediate treatment and (2) increase the likelihood that bacteria will develop resistance and will not be treatable by fidaxomicin tablets or other antibacterial drugs in the future.

Manufactured For:

Teva Pharmaceuticals

Parsippany, NJ 07054

Iss. 5/2024

PATIENT INFORMATION

Fidaxomicin (fye dax" oh mye' sin) Tablets, for oral use

What You Need to Know About Your Medicine

- Before you take fidaxomicin tablets, be sure you understand what it is for and how to take it.
- If you have questions about fidaxomicin tablets, ask your doctor or pharmacist.
- Remember that your doctor has prescribed fidaxomicin tablets only for you. Never give this medicine to anyone else.
- Keep this Patient Information for fidaxomicin tablets so you can read it again.

What are fidaxomicin tablets?

Fidaxomicin tablets are an antibiotic medicine used to treat an infection called *Clostridioides difficile*-associated diarrhea (CDAD) in adults. *Clostridioides difficile* (C-diff)

is a bacterium that can cause an infection that can damage your colon and cause stomach pain and severe diarrhea.

- Fidaxomicin tablets are not to be used to treat other types of infections in the body.
- Sometimes infections are caused by viruses rather than bacteria. Antibiotic medicines, including fidaxomicin tablets, do not kill viruses.

It is not known if fidaxomicin tablets are safe and effective in children under 6 months old.

Who should not take fidaxomicin tablets?

Do not take fidaxomicin tablets if you are allergic to fidaxomicin, or any other ingredient in fidaxomicin tablets. See the end of this Patient Information for a complete list of ingredients in fidaxomicin tablets.

What should I tell my doctor before taking fidaxomicin tablets?

Pregnancy

- If you are pregnant or plan to become pregnant, tell your doctor before you take fidaxomicin tablets.
- It is not known if fidaxomicin tablets will harm your baby while you are pregnant.
- If you are pregnant, you and your doctor should decide together if you will take fidaxomicin tablets.

Breastfeeding

- If you are breastfeeding or plan to breastfeed, tell your doctor before you take fidaxomicin tablets.
- It is not known if fidaxomicin passes into breast milk.
- If you are breastfeeding, you and your doctor should decide together if you will take fidaxomicin tablets.

Other Medicines

- **Tell your doctor about all of the medicines you take**, including prescription and over-the-counter medicines, vitamins, and herbal and dietary supplements.
- Know the medicines you take. Keep a list of your medicines to show your doctor and pharmacist when you get a new medicine.

Allergic Reactions

- See **“Who should not take fidaxomicin tablets?”**
- If you are allergic to other kinds of antibiotics called macrolides (for example: azithromycin (Zithromax®) or clarithromycin (Biaxin®)) or any other ingredient in fidaxomicin tablets, tell your doctor. See the end of this Patient Information for a complete list of ingredients in fidaxomicin tablets.

How do I take fidaxomicin tablets?

- Take fidaxomicin tablets exactly as prescribed by your doctor.
- Take fidaxomicin tablets twice a day (approximately every 12 hours). For example, if you take your first dose at 8:00 a.m. you should take your second dose at 8:00 p.m.
- You can take fidaxomicin tablets with or without food.
- **Do not** skip any doses or stop taking fidaxomicin tablets until you finish your prescribed treatment, even if you begin to feel better, unless you have a serious allergic reaction (see **“What are the possible side effects of fidaxomicin tablets?”**).

This will lower the chance that the bacteria will become resistant to fidaxomicin tablets. If this happens, fidaxomicin tablets and other antibiotic medicines may not work in the future.

What are the possible side effects of fidaxomicin tablets?

Fidaxomicin tablets can cause serious side effects, including:

- **Allergic reaction.** If you get a severe allergic reaction while taking fidaxomicin tablets, including problems breathing or shortness of breath, rash, itching or hives, or swelling of the mouth, throat, or face, stop taking fidaxomicin tablets and get emergency medical help right away.

Common side effects of fidaxomicin tablets include:

The most common side effects of fidaxomicin tablets in **adults** include:

-
- nausea
 - vomiting
 - stomach pain
 - bleeding in the stomach or intestines
 - low red blood cell count (anemia)
 - low white blood cell count

Other less common side effects of fidaxomicin tablets may include:

- swelling of any body part (such as your face, lips, tongue or around your eyes)
- itching
- hives
- bloating
- stomach tenderness
- heartburn
- problems swallowing
- high blood sugar (hyperglycemia)
- abnormal liver tests
- low levels of blood bicarbonate
- passing gas
- intestinal blockage
- serious bowel inflammation (toxic megacolon)
- low platelet count (important for clotting and to control bleeding)
- high levels of acid in your blood (metabolic acidosis)

If you have any side effect that bothers you or does not go away, tell your doctor.

There may be other side effects to fidaxomicin tablets that are not listed. Call your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.

How should I store fidaxomicin tablets?

- Store fidaxomicin tablets at room temperature between 68°F to 77°F (20°C to 25°C).
- Keep fidaxomicin tablets in its original bottle until you are ready to take it.

Keep fidaxomicin tablets and all medicines out of the reach of children.

General information about the safe and effective use of fidaxomicin tablets.

Medicines are sometimes prescribed for purposes other than those listed in the Patient Information. Do not take fidaxomicin tablets for a condition for which they were not prescribed. Do not give fidaxomicin tablets to other people, even if they have the same symptoms that you have. They may harm them. You can ask your pharmacist or doctor for information about fidaxomicin tablets that is written for health professionals.

What if I have questions?

- Call your doctor.
- Call Teva at 1-888-838-2872.

What are the ingredients in fidaxomicin tablets?

Active ingredient: fidaxomicin.

Inactive ingredients: butylated hydroxytoluene, hydroxypropyl cellulose, lecithin (soy), magnesium stearate, microcrystalline cellulose, polyethylene glycol 3350, polyvinyl alcohol-part hydrolyzed, pregelatinized corn starch, sodium starch glycolate Type A, talc, and titanium dioxide.

Pediatric use information is approved for Cubist Pharmaceuticals LLC's DIFICID® (fidaxomicin) tablets. However, due to Cubist Pharmaceuticals LLC's marketing exclusivity rights, this drug product is not labeled with that information.

Brands listed are the trademarks of their respective owners.

This Patient Information has been approved by the U.S. Food and Drug Administration.

Manufactured For:

Teva Pharmaceuticals
Parsippany, NJ 07054

Iss. 5/2024

Principal Display Panel

NDC 0480-2596-34
Fidaxomicin Tablets
200 mg
Rx only
20 Tablets

NDC 0480-2596-34

Fidaxomicin Tablets

200 mg

Rx only
20 Tablets



Each Film-Coated Tablet Contains:
Fidaxomicin 200 mg

Usual Dosage: See package insert.

Store at 20° to 25°C (68° to 77°F); excursions permitted to 15° to 30°C (59° to 86°F) [see USP Controlled Room Temperature].

Store in the original bottle.
Store out of the reach of children.

Manufactured For:
Teva Pharmaceuticals
Parsippany, NJ 07054
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52-0841

GTIN 00304802596342

0480-2596-34 2

Lot/Exp/Serialization Coding Area

FIDAXOMICIN				
fidaxomicin tablet, film coated				
Product Information				
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:0480-2596	
Route of Administration	ORAL			
Active Ingredient/Active Moiety				
	Ingredient Name	Basis of Strength	Strength	
	FIDAXOMICIN (UNII: Z5N076G8YQ) (FIDAXOMICIN - UNII:Z5N076G8YQ)	FIDAXOMICIN	200 mg	
Inactive Ingredients				
	Ingredient Name	Strength		
	BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)			
	HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH)			
	LECITHIN, SOYBEAN (UNII: 1DI56QDM62)			
	MAGNESIUM STEARATE (UNII: 70097M6I30)			
	MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)			
	POLYETHYLENE GLYCOL 3350 (UNII: G2M7P15E5P)			
	POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)			
	STARCH, CORN (UNII: O8232NY3S)			
	SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)			
	TALC (UNII: 75EV7J4R1U)			
	TITANIUM DIOXIDE (UNII: 15FIX9V2JP)			
Product Characteristics				
Color	white	Score	no score	
Shape	OVAL (modified capsule)	Size	14mm	
Flavor		Imprint Code	A205	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0480-2596-34	20 in 1 BOTTLE; Type 0: Not a Combination Product	07/15/2025	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
ANDA	ANDA208443	07/15/2025		

Labeler - Teva Pharmaceuticals, Inc. (022629579)