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## HIGHLIGHTS OF PRESCRIBING INFORMATION

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

**VAXCHORA® (Cholera Vaccine, Live, Oral)**  
**Suspension for Oral Administration**

**Initial U.S. Approval: 2016**

### RECENT MAJOR CHANGES

DOSAGE AND ADMINISTRATION, Preparation, Reconstitution and Administration (2.3)      ##/2019

### INDICATIONS AND USAGE

VAXCHORA is a vaccine indicated for active immunization against disease caused by *Vibrio cholerae* serogroup O1. VAXCHORA is approved for use in adults 18 through 64 years of age traveling to cholera-affected areas. (1)

Limitations of Use:

- The effectiveness of VAXCHORA has not been established in persons living in cholera-affected areas. (1.1)
- The effectiveness of VAXCHORA has not been established in persons who have pre-existing immunity due to previous exposure to *V. cholerae* or receipt of a cholera vaccine. (1.1)
- VAXCHORA has not been shown to protect against disease caused by *V. cholerae* serogroup O139 or other non-O1 serogroups. (1.1)

### DOSAGE AND ADMINISTRATION

- For oral administration only.
- Prepare and administer VAXCHORA in a healthcare setting equipped to dispose of medical waste. (2.3)
- Prepare VAXCHORA by reconstituting the buffer component in 100 milliliters (mL) of purified bottled or spring bottled water; then add the active component (lyophilized *V. cholerae* CVD 103-HgR). (2.3) After preparation, a single dose of VAXCHORA is 100 mL. (3)
- Instruct recipients to avoid eating or drinking for 60 minutes before and after oral ingestion of VAXCHORA. (2.2)
- Administer VAXCHORA a minimum of 10 days before potential exposure to cholera. (2.1)

component and a packet of the active component (lyophilized *V. cholerae* CVD 103-HgR). After preparation, a single dose of VAXCHORA is 100 mL. (3)

### CONTRAINDICATIONS

Severe allergic reaction (e.g., anaphylaxis) to any ingredient of VAXCHORA or to a previous dose of any cholera vaccine. (4)

### WARNINGS AND PRECAUTIONS

- The safety and effectiveness of VAXCHORA have not been established in immunocompromised persons. (5.1)
- VAXCHORA may be shed in the stool of recipients for at least 7 days. There is a potential for transmission of the vaccine strain to non-vaccinated close contacts (e.g., household contacts). Use caution when considering whether to administer VAXCHORA to individuals with immunocompromised close contacts. (5.2)

### ADVERSE REACTIONS

The most common adverse reactions (incidence > 3%) were tiredness (31%), headache (29%), abdominal pain (19%), nausea/vomiting (18%), lack of appetite (17%) and diarrhea (4%). (6)

**To report SUSPECTED ADVERSE REACTIONS, contact PaxVax, Inc. at 1-800-533-5899 or safety@paxvax.com or VAERS at 1-800-222-7967 or www.vaers.hhs.gov.**

### DRUG INTERACTIONS

Avoid concomitant administration of VAXCHORA with systemic antibiotics since these agents may be active against the vaccine strain. Do not administer VAXCHORA to patients who have received oral or parenteral antibiotics within 14 days prior to vaccination. (7.2)

Immune responses to VAXCHORA may be diminished when VAXCHORA is administered concomitantly with chloroquine. Administer VAXCHORA at least 10 days before beginning antimalarial prophylaxis with chloroquine. (7.2)

### USE IN SPECIFIC POPULATIONS

Pregnancy Registry: available at 1-800-533-5899. (8.1)

See 17 for PATIENT COUNSELING INFORMATION.

### DOSAGE FORMS AND STRENGTHS

Suspension for oral administration supplied as a packet of the buffer

Revised: ##/2019

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## FULL PRESCRIBING INFORMATION

### 1 INDICATIONS AND USAGE

VAXCHORA is a vaccine indicated for active immunization against disease caused by *Vibrio cholerae* serogroup O1 in adults 18 through 64 years of age traveling to cholera-affected areas.

#### 1.1 Limitations of Use

The effectiveness of VAXCHORA has not been established in persons living in cholera-affected areas.

The effectiveness of VAXCHORA has not been established in persons who have pre-existing immunity due to previous exposure to *V. cholerae* or receipt of a cholera vaccine.

VAXCHORA has not been shown to protect against disease caused by *V. cholerae* serogroup O139 or other non-O1 serogroups.

### 2 DOSAGE AND ADMINISTRATION

For oral administration only.

#### 2.1 Dose and Schedule

Administer a single oral dose of VAXCHORA a minimum of 10 days before potential exposure to cholera.

The safety and effectiveness of revaccination with VAXCHORA have not been established.

#### 2.2 Restrictions on Eating and Drinking

Instruct recipients to avoid eating or drinking for 60 minutes before and after oral ingestion of VAXCHORA.

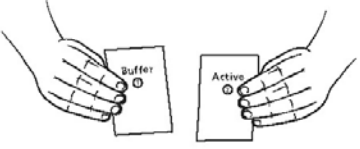

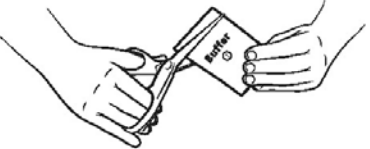


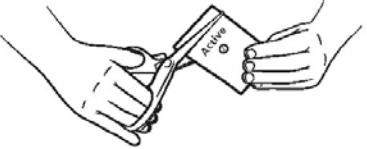


#### 2.3 Preparation, Reconstitution and Administration

Prepare and administer VAXCHORA in a healthcare setting equipped to dispose of medical waste [see *Disposal Instructions (2.4)*].

1. Reconstitution should be completed within 15 minutes of removing the carton from the refrigerator [see *Storage and Handling (16.2)*]. Locate the 2 packets: the buffer component (Packet 1) and the active component (Packet 2).

2. Pour 100 mL of cold or room temperature (41°F-72°F; 5°C-22°C) purified bottled or spring bottled water into a clean, disposable cup. Do not use tap water, sparkling (carbonated) water, non-purified or non-spring bottled water, other beverages, or other liquids.
3. Use scissors to cut the top off the buffer component packet.
4. Empty buffer component packet contents into cup. Effervescence will occur.
5. Using a disposable stirrer, stir until the buffer component completely dissolves.
6. Use scissors to cut the top off the active component packet.
7. Empty the active component packet contents (lyophilized *V. cholerae* CVD 103-HgR) into the cup containing the buffer solution.
8. Stir for at least 30 seconds and until active component disperses to form a slightly cloudy suspension that may contain some white particulates. The active component may not dissolve completely.
9. VAXCHORA must be consumed within 15 minutes of reconstitution. The recipient should drink the full contents of the cup at once. Some residue may remain in the cup and should be discarded with the cup.

**NOTE:** If the packets are reconstituted in the improper order, the vaccine must be discarded [*see Disposal Instructions (2.4)*].

<p>1</p>  <p>Reconstitution should be completed within 15 minutes of removing the carton from the refrigerator [see <i>Storage and Handling</i> (16.2)]. Locate the 2 packets: the buffer component (Packet 1) and the active component (Packet 2).</p>	<p>2</p>  <p>Pour 100 mL of cold or room temperature (41°F-72°F; 5°C-22°C) purified bottled or spring bottled water into a clean, disposable cup. Do not use tap water, sparkling (carbonated) water, non-purified or non-spring bottled water, other beverages, or other liquids.</p>	<p>3</p>  <p>Use scissors to cut the top off the buffer component packet.</p>
<p>4</p>  <p>Empty buffer component packet contents into cup. Effervescence will occur.</p>	<p>5</p>  <p>Using a disposable stirrer, stir until the buffer component completely dissolves.</p>	<p>6</p>  <p>Use scissors to cut the top off the active component packet.</p>
<p>7</p>  <p>Empty the active component packet contents (lyophilized <i>V. cholerae</i> CVD 103-HgR) into the cup containing the buffer solution.</p>	<p>8</p>  <p>Stir for at least 30 seconds and until active component disperses to form a slightly cloudy suspension that may contain some white particulates. The active component may not dissolve completely.</p>	<p>9</p>  <p>VAXCHORA must be consumed within 15 minutes of reconstitution. The recipient should drink the full contents of the cup at once. Some residue may remain in the cup and should be discarded with the cup.</p>

## **2.4 Disposal Instructions**

Dispose of the cup, packets and stirrer according to standard procedures for medical waste.

Inactivate any spilled vaccine and clean any non-disposable equipment used in the preparation of VAXCHORA with 70% isopropyl alcohol or 10% bleach solution.

## **3 DOSAGE FORMS AND STRENGTHS**

VAXCHORA is a suspension for oral administration. Before reconstitution, each dose of VAXCHORA is supplied as a foil packet of buffer and an accompanying foil packet of the active component (lyophilized *V. cholerae* CVD 103-HgR). After reconstitution, a single dose of VAXCHORA is 100 milliliters (mL).

## **4 CONTRAINDICATIONS**

Do not use in persons who have a history of severe allergic reaction (e.g., anaphylaxis) to any ingredient of VAXCHORA or to a previous dose of any cholera vaccine [*see Description (11)*].

## **5 WARNINGS AND PRECAUTIONS**

### **5.1 Altered Immunocompetence**

The safety and effectiveness of VAXCHORA have not been established in immunocompromised persons [*see Immunocompromised Individuals (8.6)*].

### **5.2 Shedding and Transmission**

VAXCHORA may be shed in the stool of recipients for at least 7 days. There is a potential for transmission of the vaccine strain to non-vaccinated close contacts (e.g., household contacts) [*see Pharmacodynamics (12.2)*]. Use caution when considering whether to administer VAXCHORA to individuals with immunocompromised close contacts.

## **6 ADVERSE REACTIONS**

### **6.1 Clinical Trials Experience**

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

The safety of VAXCHORA was evaluated in four randomized, placebo-controlled, multicenter clinical trials. A total of 3235 adults 18 through 64 years of age received one dose of VAXCHORA and 562 received placebo [physiologic saline (N=551) or lactose (N=11)]. Overall, the mean age was 32.5 years; 53.8% of trial participants were female; 67.1% were White, 27.3% were Black or African American, 1.8% were Asian, 1.7% were multiracial, 1.3% were other, 0.6% were American Indian or Alaskan Native and 0.3% were Native Hawaiian or Pacific Islander. There were 9.3% Hispanic or Latino participants.

### Solicited Adverse Reactions

Adults 18 through 45 years of age received VAXCHORA in a multi-center, double-blind, randomized (8:1), placebo-controlled trial conducted in the United States and Australia (Study 1). The safety analysis set included 2789 VAXCHORA recipients. Solicited adverse reactions were recorded daily for 7 days following vaccination. Table 1 presents the frequency and severity of solicited adverse reactions observed within 7 days following receipt of VAXCHORA or placebo in Study 1.

**Table 1: Rates of Solicited Adverse Reactions Reported in VAXCHORA Trial Participants 18 to 45 Years of Age During 7 Days Post-Vaccination**

Adverse Reaction	Study 1 <sup>a</sup>	
	VAXCHORA (N=2789) <sup>b</sup>	Placebo (Saline) (N=350) <sup>b</sup>
	%	%
Tiredness	31.3	27.4
Mild	18.7	16.3
Moderate	12.0	9.9
Severe <sup>c</sup>	0.7	1.2
Headache	28.9	23.6
Mild	18.9	14.6
Moderate	9.6	8.8
Severe <sup>c</sup>	0.5	0.3
Abdominal Pain	18.7	16.9
Mild	12.1	12.0
Moderate	6.2	5.0
Severe <sup>c</sup>	0.4	0.0
Nausea/Vomiting	18.3	15.2
Mild	13.3	11.4
Moderate	4.7	3.8
Severe <sup>c</sup>	0.3	0.0
Lack of Appetite	16.5	16.6
Mild	11.7	12.2
Moderate	4.4	4.4

Severe <sup>c</sup>	0.3	0.0
Diarrhea	3.9	1.2
Mild	2.4	0.9
Moderate	0.7	0.3
Severe <sup>c</sup>	0.84	0.0
Fever	0.6	1.2
Mild	0.2	0.3
Moderate	0.3	0.9
Severe <sup>c</sup>	0.11	0.0

<sup>a</sup>Data are derived from Study 1 (NCT02094586).

<sup>b</sup>N represents number of subjects who completed a memory aid.

<sup>c</sup> Severe category includes both grade 3 (severe) and grade 4 (potentially life-threatening) adverse events.

Grading scales are defined as follows:

Tiredness, Headache, Abdominal Pain, Nausea, Lack of Appetite: Mild = no interference with activity, Moderate = Some interference with activity, Severe = significant, prevents daily activity, Potentially Life Threatening = emergency room (ER) visit or hospitalization.

Vomiting: Mild = 1-2 episodes/24 hours, Moderate = >2 episodes/24 hours, Severe = requires intravenous hydration, Potentially Life Threatening = ER visit or hospitalization for hypotensive shock.

Diarrhea: Mild = 4 loose stools/24 hours, Moderate = 5 loose stools/24 hours, Severe =  $\geq 6$  loose stools /24 hours, Potentially Life Threatening = ER visit or hospitalization.

Fever: Mild = 38.0-38.4°C/100.4-101.1°F, Moderate = 38.5-38.9°C/101.2-102.0°F, Severe = 39.0-40.0°C/102.1-104.0°F, Potentially Life Threatening = >40.0 °C/104.0 °F.

## Serious Adverse Events

In a pooled analysis of the four clinical studies, 0.6% (20/3235) of VAXCHORA recipients and 0.5% (3/562) of placebo recipients reported a serious adverse event within 6 months post-vaccination. None of these events were considered to be related to vaccination.

## 7 DRUG INTERACTIONS

### 7.1 Food and Drink

Avoid food or drink for 60 minutes before and after vaccine administration [*see Restrictions on Eating and Drinking (2.2)*].

### 7.2 Concomitant Vaccines or Medications

#### Vaccines

No data are available on concomitant administration of VAXCHORA with other vaccines.



## Antibiotics

Avoid concomitant administration of VAXCHORA with systemic antibiotics since these agents may be active against the vaccine strain and prevent a sufficient degree of multiplication to occur in order to induce a protective immune response. Do not administer VAXCHORA to patients who have received oral or parenteral antibiotics within 14 days prior to vaccination.

## Antimalarial Prophylaxis

Data from a study with a similar product indicate that the immune responses to VAXCHORA may be diminished when VAXCHORA is administered concomitantly with chloroquine. Administer VAXCHORA at least 10 days before beginning antimalarial prophylaxis with chloroquine.

### **7.3 Immunosuppressive Treatments**

Immunosuppressive therapies, including irradiation, antimetabolites, alkylating agents, cytotoxic drugs and corticosteroids (used in greater than physiologic doses), may reduce the immune response to VAXCHORA [*see Use in Specific Populations (8.6)*].

## **8 USE IN SPECIFIC POPULATIONS**

### **8.1 Pregnancy**

#### Pregnancy Exposure Registry

There is a pregnancy exposure registry that monitors pregnancy outcomes in women exposed to VAXCHORA during pregnancy. To enroll in or obtain information about the registry, please call PaxVax at 1-800-533-5899.

#### Risk Summary

VAXCHORA is not absorbed systemically following oral administration, and maternal use is not expected to result in fetal exposure to the drug.

#### Clinical Considerations

##### *Disease-associated maternal and/or embryo/fetal risk*

Maternal cholera disease is associated with adverse pregnancy outcomes including fetal death.

##### *Fetal/neonatal adverse reactions*

The vaccine strain may be shed in the stool of the vaccinated mother for at least 7 days, with a potential for transmission of the vaccine strain from mother to infant during vaginal delivery.

### **8.2 Lactation**

## Risk Summary

VAXCHORA is not absorbed systemically by the mother following oral administration, and breastfeeding is not expected to result in exposure of the child to VAXCHORA.

### **8.4 Pediatric Use**

The safety and effectiveness of VAXCHORA have not been established in children and adolescents younger than 18 years.

### **8.5 Geriatric Use**

The safety and effectiveness of VAXCHORA have not been established in adults 65 years of age or older.

### **8.6 Immunocompromised Individuals**

The safety and effectiveness of VAXCHORA have not been established in immunocompromised individuals. The immunologic response to VAXCHORA may be diminished in immunocompromised individuals [*see Drug Interactions (7.3)*].

## **11 DESCRIPTION**

VAXCHORA (Cholera Vaccine, Live, Oral) is a live, attenuated bacterial vaccine suspension for oral administration containing the *V. cholerae* strain CVD 103-HgR. CVD 103-HgR was constructed from the serogroup O1 classical Inaba strain 569B by deleting the catalytic domain sequence of both copies of the *ctxA* gene, which prevents the synthesis of active cholera toxin (CT). This attenuated strain remains able to synthesize the immunogenic non-toxic B subunit of CT (encoded by the *ctxB* gene). In addition, a marker was inserted into the hemolysin gene locus (*hlyA*) to enable differentiation of the vaccine strain from wild type *V. cholerae* O1.

The vaccine strain is grown in fermentors under controlled conditions in medium containing casamino acids, yeast extract, mineral salts, and an anti-foaming agent. The bacteria are concentrated by ultrafiltration before addition of a stabilization solution containing ascorbic acid (an antioxidant), Hy-Case SF (hydrolyzed casein [a protein derived from cow's milk], a cryoprotectant), and sucrose (a cryoprotectant). The stabilized bacteria are lyophilized, milled, and blended with anhydrous lactose (a bulking agent). The active component blend is filled into packets.

The buffer component is manufactured by blending together sodium bicarbonate (a gastric acid neutralizer), sodium carbonate (a buffer), ascorbic acid (a buffer and water chlorine neutralizer), and dried lactose (a manufacturing flow aid). The buffer component blend is filled into packets. One buffer component packet and one active component packet are packaged into individual single dose cartons for distribution.

After reconstitution, VAXCHORA contains  $4 \times 10^8$  to  $2 \times 10^9$  colony forming units (CFU) of live attenuated *V. cholerae* CVD 103-HgR. The resulting suspension should be slightly cloudy and may contain white particulates. The active and buffer ingredients are shown in Table 2.

**Table 2: Vaccine Composition**

Ingredient	Quantity/packet
Active Component Packet	
<i>V. cholerae</i> CVD 103-HgR	$4 \times 10^8$ to $2 \times 10^9$ CFU <sup>a</sup>
Sucrose	$\leq 165.37$ mg <sup>b</sup>
Hy-Case SF (hydrolyzed casein)	$\leq 17.11$ mg
Ascorbic acid	$\leq 8.55$ mg
Anhydrous lactose	$\leq 2.09$ g <sup>c</sup>
Buffer Component Packet	
Sodium bicarbonate	2.16–2.41 g
Sodium carbonate	0.24–0.49 g
Ascorbic acid	1.50–1.80 g
Dried lactose	0.18–0.22 g

<sup>a</sup> CFU=colony forming units.

<sup>b</sup> mg=milligrams.

<sup>c</sup> g=grams.

## 12 CLINICAL PHARMACOLOGY

### 12.1 Mechanism of Action

VAXCHORA contains live attenuated cholera bacteria that replicate in the gastrointestinal tract of the recipient. Immune mechanisms conferring protection against cholera following receipt of VAXCHORA have not been determined. However, rises in serum vibriocidal antibody 10 days after vaccination with VAXCHORA were associated with protection in a human challenge study (Study 2) [See Immunogenicity (14.2)].

### 12.2 Pharmacodynamics

Shedding of the vaccine strain was evaluated in the first 7 days post-vaccination in a study of 53 healthy adult vaccine recipients (Study 3). VAXCHORA was shed in the stools of 11.3% [95% CI 4.3%, 23.0%] of vaccine recipients on any day through 7 days post-vaccination. During the 7 days post-vaccination, the proportion of subjects shedding was highest on day 7 (7.5% [95% CI 2.1%, 18.2%]). The duration of shedding of the vaccine strain is unknown.

## 13 NONCLINICAL TOXICOLOGY

### 13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

VAXCHORA has not been evaluated for the potential to cause carcinogenicity or genotoxicity, or to impair fertility.

## 14 CLINICAL STUDIES

### 14.1 Efficacy Against *V. cholerae* Challenge

Study 2 was a randomized, double-blind, saline placebo-controlled *V. cholerae* challenge study conducted in the US. Subjects 18 through 45 years of age (N=197) with no prior history of cholera infection or travel to a cholera-endemic area in the previous 5 years were randomized according to a 1:1 ratio to receive one dose of VAXCHORA or placebo. In order to identify the subset of subjects to be challenged, an unblinded statistician prepared four randomly ordered lists of subjects per site, one list each for vaccine recipients with blood type O, vaccine recipients with non-O blood types, placebo recipients with blood type O, and placebo recipients with non-O blood types. This was done to maintain a minimum of 60% blood group O subjects in each treatment group. Individuals with type O blood are less likely to be infected with *V. cholerae*, but are at risk for developing severe cholera if infected. Each site was provided with a blinded version of the four lists specific to its site and advised on the number of subjects from each list to challenge. In the event that a subject was determined to be ineligible for challenge, the site was instructed to select the next subject from the same list as the ineligible subject.

The challenges were split into 2 cohorts for 10 day and 3 month challenges. Subjects were admitted to an inpatient unit. Subjects had nothing by mouth from midnight before ingestion of the challenge strain, except for water, and had nothing by mouth for 90 minutes after ingestion of the challenge strain. Approximately 1 minute prior to challenge, subjects ingested 120 mL sodium bicarbonate (NaHCO<sub>3</sub>) buffer. The oral challenge consisted of 1 x 10<sup>5</sup> CFU live wild type *V. cholerae* El Tor Inaba N16961 in 30 mL NaHCO<sub>3</sub> buffer at 10 days or 3 months post-vaccination. The co-primary objectives were to demonstrate the efficacy of a single dose of VAXCHORA in the prevention of moderate to severe diarrhea following challenge at 10 days and 3 months post-vaccination. Moderate to severe diarrhea was defined as cumulative diarrheal purge  $\geq$  3 liters (L) within 10 days after challenge. Diarrheal stool was defined as  $\geq$  2 unformed stools (takes shape of container) collected during a 48 hour period  $\geq$  200 grams (g) or a single unformed stool  $\geq$  300 g. Subjects were instructed to collect every stool from the time of challenge until discharge from the inpatient unit. Nursing staff or study personnel inspected all stool, graded the consistency of the stool and calculated the total weight of diarrheal stool per day. Weight of stool was converted to volume using the formula 1 g=1 mL. VAXCHORA recipients challenged at 10 days post-vaccination and VAXCHORA recipients challenged at 3 months post-vaccination were compared with a pooled group of placebo (saline) recipients challenged at 10 days or 3 months post-vaccination.

Of the 95 VAXCHORA recipients, 68 were challenged; 35 were challenged at 10 days post-vaccination and 33 were challenged at 3 months post-vaccination. Of the 102 placebo recipients, 66 were challenged; 33 were challenged at 10 days post-vaccination and 33 at 3 months post-vaccination. Among all randomized subjects, the mean age was 31.0 years. Overall, the mean age of the challenge population was 31.4 years. More males were in the vaccine group (71.6%) compared to the placebo group (54.9%). The majority of randomized subjects were Black (67.5%), 29.4% were White, 0.5% were American Indian/Alaskan Native, 0.5% were Asian, and 2.0% were other. There were 4.6% Hispanic or Latino participants. Overall 50.3% had blood type O. Among subjects selected for either challenge cohort, more males were challenged in the vaccine group (76.5%) compared to the placebo group (57.6%). The majority (70.9%) of the challenge population were Black, 25.4% were White, 0.7% were American Indian/Alaskan Native, 0.7% were Asian, and 2.2% were other. There were 3.7% Hispanic or Latino participants. Overall, 56.0% of challenged subjects had blood type O.

Vaccine efficacy against the occurrence of moderate to severe diarrhea at 10 days post-vaccination was 90.3% [95% CI 62.7%, 100.0%] and at 3 months post-vaccination was 79.5% [95% CI 49.9%, 100.0%] (Table 3).

**Table 3: Vaccine Efficacy in the Prevention of Moderate to Severe Diarrhea Following Challenge with *V. cholerae* O1 El Tor Inaba at 10 Days and 3 Months Post-Vaccination (Intent-to-Treat Population)**

Parameter	VAXCHORA 10 Day Challenge <sup>a,c</sup> N=35 <sup>d</sup>	VAXCHORA 3 Month Challenge <sup>a,c</sup> N=33 <sup>d</sup>	Combined Placebo <sup>a,b</sup> 10 Day or 3 Month Challenge <sup>c</sup> N=66 <sup>d</sup>
Number of Subjects with Moderate or Severe Diarrhea (Attack Rate) <sup>e</sup>	2 (5.7%)	4 (12.1%)	39 (59.1%)
Vaccine Efficacy % <sup>f,g</sup> [95% CI <sup>h</sup> ]	90.3% [62.7%, 100.0%]	79.5% [49.9%, 100.0%]	

<sup>a</sup> Data are derived from Study 2 (NCT01895855).

<sup>b</sup> Combined placebo group comprised of all placebo recipients who were challenged at either 10 days (N=33) or 3 months (N=33) following vaccination.

<sup>c</sup> Challenge strain was *V. cholerae* O1 El Tor Inaba N16961.

<sup>d</sup> N=number of subjects challenged in each group.

<sup>e</sup> Moderate or severe diarrhea ( $\geq 3$  liters of diarrhea) within 10 days after challenge.

<sup>f</sup> Vaccine Efficacy= $[(\text{Attack Rate in Placebo Group} - \text{Attack Rate in Vaccine Group})/\text{Attack Rate in Placebo Group}] \times 100$ .

<sup>g</sup> Pre-specified criteria for success were that the lower bound of the two-sided 95% confidence interval for vaccine efficacy must be  $\geq 30\%$  in both the 10 Day and 3 Month challenge groups.

<sup>h</sup> CI=confidence interval.

## 14.2 Immunogenicity

### *Vibriocidal Antibody Against the Vaccine Strain (classical Inaba)*

A vibriocidal antibody assay was used to measure serum levels of neutralizing antibodies against the vaccine strain.

Study 2 was a randomized, double-blind, saline placebo-controlled *V. cholerae* challenge study conducted in adults 18 through 45 years of age. In the subset of subjects challenged in Study 2, 91% [95% CI 82%, 97%] of vaccinees seroconverted prior to challenge and 9% developed moderate to severe cholera following challenge, while 2% of placebo recipients seroconverted prior to challenge and 59% developed moderate to severe cholera following challenge. (Seroconversion was defined as a  $\geq 4$ -fold rise in serum vibriocidal antibody from baseline to 10 days post-vaccination.) Based on the observed association between seroconversion and protection from *V. cholerae* disease, seroconversion rate at 10 days post-vaccination was used to evaluate response to vaccination in other age groups.

Study 1 was a randomized, double-blind, saline placebo-controlled safety and immunogenicity study conducted in the US and Australia. A total of 3146 subjects 18 through 45 years of age not previously exposed to cholera were randomized 8:1 to receive one dose of VAXCHORA or placebo. The mean age was 29.9 years; 45.2% were male; 68.3% were White, 25.6% were Black, 2.0% were Asian, 1.9% were multiracial, 1.4% were other, 0.4% were American Indian/Alaskan Native, and 0.3% were Native Hawaiian/Pacific Islander. There were 10.0% Hispanic or Latino participants.

In this study, the rates of seroconversion were 93.5% [95% CI 92.5%, 94.4%] in vaccine recipients and 4% [95% CI 2%, 7%] in placebo recipients at 10 days post-vaccination.

Study 4 was a randomized, double-blind, placebo-controlled safety and immunogenicity study conducted in the US. A total of 398 subjects 46 through 64 years of age with no prior history of cholera infection or travel to a cholera-endemic area in the previous 5 years were randomized 3:1 to receive one dose of VAXCHORA or placebo. Overall, the mean age of the randomized population was 53.8 years; 45.7% were male; 74.9% were White, 21.9% were Black, 1.8% were American Indian/Alaskan Native, 0.5% were Asian, 0.5% were multiracial, 0.3% were Native Hawaiian/Pacific Islander, and 0.3% were other. There were 7.5% Hispanic or Latino participants.

Seroconversion rates at 10 days post-vaccination by classical Inaba vibriocidal antibody among 46 through 64 year old subjects in Study 4 were compared to those in 18 through 45 year old subjects in Study 1. VAXCHORA recipients from Study 1 were in the same age group as those in Study 2, the *V. cholerae* challenge study.

Adults 46 through 64 years were shown to have a non-inferior rate of seroconversion by classical Inaba vibriocidal antibody at 10 days post-vaccination compared to adults 18 through 45 years of age (Table 4).

**Table 4: Vibriocidal Antibody Seroconversion Against Classical Inaba *V. cholerae* Vaccine Strain at 10 Days Post-Vaccination in Adults 46 through 64 Years of Age (Study 4) Compared to Adults 18 through 45 Years of Age (Study 1) [Bridging Analysis Population]**

Study <sup>a</sup>	Dose/CFU	VAXCHORA N <sup>b</sup>	VAXCHORA Seroconversion % [95% CI <sup>c</sup> ]
Study 4 (46 through 64 year olds)	1 x 10 <sup>9</sup>	291	90.4% [86.4%, 93.5%]
Study 1 (18 through 45 year olds)	1 x 10 <sup>9</sup>	2687	93.5% [92.5%, 94.4%]
Difference in Seroconversion Rates <sup>d,e</sup>			-3.1% [-6.7%, 0.4%]

<sup>a</sup> Data are derived from Study 1 (NCT02094586) and Study 4 (NCT02100631).

<sup>b</sup> N=number of subjects with analyzable samples at Day 1 and Day 11.

<sup>c</sup> CI=confidence interval.

<sup>d</sup> Seroconversion is defined as the percentages of subjects who had at least a 4-fold rise in vibriocidal antibody titer at 10 days post-vaccination compared to baseline.

<sup>e</sup> Pre-specified non-inferiority criterion was that the lower bound of the two-sided 95% confidence interval on the difference in seroconversion rate (Study 4 minus Study 1) must be greater than -10 percentage points.

#### *Vibriocidal Antibody Against Classical Ogawa, El Tor Inaba and El Tor Ogawa*

*V. cholerae* serogroup O1 consists of four major subtypes: classical Inaba, classical Ogawa, El Tor Inaba and El Tor Ogawa. Serum vibriocidal antibody against the three types of *V. cholerae* not contained in the vaccine, namely classical Ogawa, El Tor Inaba and El Tor Ogawa, was also measured in Study 2 and Study 4. The percentages of vaccine recipients who seroconverted against each of the 4 major biotype/serotypes of *V. cholerae* serogroup O1 at 10 days post-vaccination (71.4% to 91.4%) are shown in Table 5.

**Table 5: Seroconversion Rates 10 Days Post-Vaccination for the Four Major *V. cholerae* O1 Serogroup Biotypes and Serotypes in Studies 2 and 4 [Immunogenicity Evaluable Population]**

Cholera Strain	Study 2 <sup>a</sup> (18 through 45 year olds) VAXCHORA		Study 4 <sup>a</sup> (46 through 64 year olds) VAXCHORA	
	N <sup>b</sup>	% <sup>c</sup> [95% CI <sup>d</sup> ]	N <sup>b</sup>	% [95% CI]
Classical Inaba <sup>e</sup>	93	90.3% [82.4%, 95.5%]	291	90.4% [86.4%, 93.5%]
El Tor Inaba	93	91.4% [83.8%, 96.2%]	290	91.0% [87.1%, 94.1%]
Classical Ogawa	93	87.1% [78.5%, 93.2%]	291	73.2% [67.7%, 78.2%]
El Tor Ogawa	93	89.2% [81.1%, 94.7%]	290	71.4% [65.8%, 76.5%]

<sup>a</sup> Data are derived from Study 2 (NCT01895855) and Study 4 (NCT02100631).

<sup>b</sup> N=number of subjects with measurements at baseline and 10 days post-vaccination. One subject in Study 2 did not have a Day 11 measurement and was dropped from the analysis.

<sup>c</sup> Seroconversion is defined as the percentages of subjects who had at least a 4-fold rise in vibriocidal antibody titer at 10 days post-vaccination compared to the titer measured at baseline.

<sup>d</sup> CI=confidence interval.

<sup>e</sup> VAXCHORA contains the classical Inaba strain of *V. cholerae* O1.

## 16 HOW SUPPLIED/STORAGE AND HANDLING

### 16.1 How Supplied

VAXCHORA is supplied as shown in Table 6. The contents of both packets are reconstituted with purified bottled or spring bottled water, to form one oral dose of the vaccine.

**Table 6: VAXCHORA Product Presentation**

Presentation	Carton NDC Number	Components
Single dose carton containing two packets	NDC 70460-001-01	Buffer Component Packet NDC 70460-003-02 Active Component Packet NDC 70460-002-02

### 16.2 Storage and Handling

Store VAXCHORA buffer component and active component packets refrigerated at 36°F to 46°F (2°C to 8°C).



Protect from light and moisture.

Packets should not be out of refrigerated storage for more than 15 minutes prior to reconstitution; when out of refrigerated storage, packets should not be exposed to temperatures above 80°F (27°C).

## 17 PATIENT COUNSELING INFORMATION

Prior to administration of this vaccine, the health care professional should inform the individual of the following:

- Advise vaccine recipients to exercise caution regarding food and water consumed in cholera-affected areas, in accordance with the recommendations from the Centers for Disease Control and Prevention for the prevention of cholera in travelers.
- Educate vaccine recipients regarding the most common adverse reactions occurring within 7 days post-vaccination with VAXCHORA (tiredness, headache, abdominal pain, nausea/vomiting, lack of appetite, and diarrhea).
- Inform vaccine recipients that VAXCHORA is a live attenuated vaccine and has the potential for transmission of the vaccine strain to close contacts (e.g., household contacts). For at least 14 days following vaccination with VAXCHORA, vaccine recipients should wash their hands thoroughly after using the bathroom and before preparing or handling food.
- Register women who receive VAXCHORA while pregnant in the pregnancy registry by calling 1-800-533-5899 [*see Pregnancy (8.1)*].
- Instruct vaccine recipients to report adverse reactions to their healthcare provider.

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