

Monodox® Doxycycline Monohydrate Capsules Revised: November 2011 Rx only 102-ZZ

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Rx only

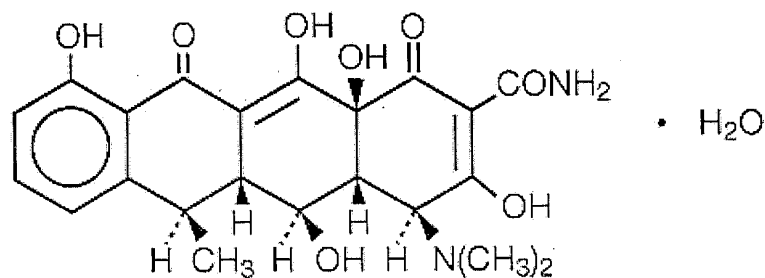
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To reduce the development of drug-resistant bacteria and maintain the effectiveness of Monodox and other antibacterial drugs, Monodox should be used only to treat or prevent infections that are proven or strongly suspected to be caused by bacteria.

DESCRIPTION

Doxycycline is a broad-spectrum antibiotic synthetically derived from oxytetracycline. Monodox 100 mg, 75 mg, and 50 mg capsules contain doxycycline monohydrate equivalent to 100 mg, 75 mg, or 50 mg of doxycycline for oral administration. The chemical designation of the light-yellow crystalline powder is alpha-6-deoxy-5-oxytetracycline.

Structural formula:



C₂₂H₂₄N₂O₈ • H₂O

M.W. = 462.45

Doxycycline has a high degree of lipid solubility and a low affinity for calcium binding. It is highly stable in normal human serum. Doxycycline will not degrade into an epianhydro form.

Inert ingredients: colloidal silicon dioxide; magnesium stearate; microcrystalline cellulose; sodium starch glycolate; and a hard gelatin capsule which contains black iron oxide, red iron oxide, titanium dioxide, and yellow iron oxide for the 100 mg and 75 mg strengths, titanium dioxide and yellow iron oxide for the 50 mg strength. The capsules are printed with edible ink containing black iron oxide, red iron oxide, and yellow iron oxide for the 50 mg and 100 mg strengths and black iron oxide, FD&C Blue No. 2, FD&C Red No. 40, FD&C Blue No. 1, and D&C Yellow No. 10 for the 75 mg strength.

CLINICAL PHARMACOLOGY

Tetracyclines are readily absorbed and are bound to plasma proteins in varying degrees. They are concentrated by the liver in the bile and excreted in the urine and feces at high concentrations in a biologically active form. Doxycycline is virtually completely absorbed after oral administration.

Following a 200 mg dose of doxycycline monohydrate, 24 normal adult volunteers averaged the following serum concentration values:

Time (hr):	0.5	1.0	1.5	2.0	3.0	4.0	8.0	12.0	24.0	48.0	72.0
Conc.	1.02	2.26	2.67	3.01	3.16	3.03	2.03	1.62	0.95	0.37	0.15 (µg/mL)

Average Observed Values

Maximum Concentration	3.61 µg/mL (± 0.9 sd)
Time of Maximum Concentration	2.60 hr (± 1.10 sd)
Elimination Rate Constant	0.049 per hr (± 0.030 sd)
Half-Life	16.33 hr (± 4.53 sd)

Excretion of doxycycline by the kidney is about 40%/72 hours in individuals with normal function (creatinine clearance about 75 mL/min). This percentage excretion may fall as low as 1-5%/72 hours in individuals with severe renal insufficiency (creatinine clearance below 10 mL/min). Studies have shown no significant difference in serum half-life of doxycycline (range 18-22 hours) in individuals with normal and severely impaired renal function.

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Hemodialysis does not alter serum half-life.

Microbiology: The tetracyclines are primarily bacteriostatic and are thought to exert their antimicrobial effect by the inhibition of protein synthesis. The tetracyclines, including doxycycline, have a similar antimicrobial spectrum of activity against a wide range of gram-positive and gram-negative microorganisms. Cross-resistance of these microorganisms to tetracyclines is common.

Doxycycline has been shown to be active against most strains of the following microorganisms, both *in vitro* and in clinical infections as described in the INDICATIONS AND USAGE section.

Aerobic Gram-Positive Microorganisms:

Because many strains of the following groups of gram-positive microorganisms have been shown to be resistant to tetracyclines, culture and susceptibility testing are recommended.

Bacillus anthracis
Listeria monocytogenes
*Staphylococcus aureus**

*Doxycycline is not the drug of choice in the treatment of any type of staphylococcal infection.

Up to 44 percent of strains of *Streptococcus pyogenes* and 74 percent of *Streptococcus faecalis* have been found to be resistant to tetracycline drugs. Therefore, tetracyclines should not be used to treat streptococcal infections unless the microorganism has been demonstrated to be susceptible.

Streptococcus pneumoniae

Aerobic Gram-Negative Microorganisms:

<i>Bartonella bacilliformis</i>	<i>Haemophilus ducreyi</i>
<i>Brucella species</i>	<i>Haemophilus influenzae</i>
<i>Calymmatobacterium granulomatis</i>	<i>Neisseria gonorrhoeae</i>
<i>Campylobacter fetus</i>	<i>Vibrio cholerae</i>
<i>Francisella tularensis</i>	<i>Yersinia pestis</i>

Because many strains of the following groups of gram-negative microorganisms have been shown to be resistant to tetracyclines, culture and susceptibility testing are recommended:

<i>Acinetobacter species</i>	<i>Klebsiella species</i>
<i>Enterobacter aerogenes</i>	<i>Shigella species</i>
<i>Escherichia coli</i>	

Anaerobic Microorganisms:

<i>Actinomyces israelii</i>	<i>Fusobacterium fusiforme</i>
<i>Clostridium species</i>	

Other Microorganisms:

<i>Borrelia recurrentis</i>	<i>Rickettsiae</i>
<i>Chlamydia psittaci</i>	<i>Treponema pallidum</i>
<i>Chlamydia trachomatis</i>	<i>Treponema pertenue</i>
<i>Mycoplasma pneumoniae</i>	

Susceptibility Tests:

Dilution Techniques:

Quantitative methods are used to determine antimicrobial minimum inhibitory concentrations (MIC's). These MIC's provide estimates of the susceptibility of bacteria to antimicrobial compounds. The MIC's should be determined using a standardized procedure. Standardized procedures are based on a dilution method^{1,3} (broth or agar) or equivalent with standardized inoculum concentrations and standardized concentrations of tetracycline powder. The MIC values should be interpreted according to the following criteria:

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Interpretive criteria for *Enterobacteriaceae*, *Staphylococcus aureus* and *Acinetobacter* spp.

Microorganism	MIC Interpretive Standard (µg/mL)		
	Susceptible (S)	Intermediate (I)	Resistant (R)
<i>Enterobacteriaceae</i>	≤ 4	8	≥ 16
<i>Staphylococcus aureus</i>			
<i>Acinetobacter</i> spp.			

Microorganisms that are susceptible to tetracycline are generally susceptible to doxycycline.

Interpretive Criteria for *Haemophilus* spp.

Microorganism	MIC Interpretive Standard (µg/mL)		
	Susceptible (S)	Intermediate (I)	Resistant (R)
<i>Haemophilus</i> spp.	≤ 2	4	≥ 8

Interpretive criteria for *Haemophilus* spp. are applicable only to tests performed by broth microdilution method using *Haemophilus* Test Medium (HTM).^{1,3}

Microorganisms that are susceptible to tetracycline are generally susceptible to doxycycline.

Interpretive Criteria for *Neisseria gonorrhoeae*

Microorganism	MIC Interpretive Standard (µg/mL)		
	Susceptible (S)	Intermediate (I)	Resistant (R)
<i>Neisseria gonorrhoeae</i>	≤ 0.25	0.5 - 1	≥ 2

Interpretive criteria for *Neisseria gonorrhoeae* are applicable only to tests performed by agar dilution method using GC agar base with 1% defined growth supplement.^{1,3}

Microorganisms that are susceptible to tetracycline are generally susceptible to doxycycline.

Interpretive Criteria for *Streptococcus pneumoniae*

Microorganism	MIC Interpretive Standard (µg/mL)		
	Susceptible (S)	Intermediate (I)	Resistant (R)
<i>Streptococcus pneumoniae</i>	≤ 2	4	≥ 8

Interpretive criteria for *Streptococcus pneumoniae* are applicable only to tests performed by broth microdilution method using Cation-Adjusted Mueller-Hinton broth with 2.5% - 5% lysed horse blood.^{1,3}

Microorganisms that are susceptible to tetracycline are generally susceptible to doxycycline.

Interpretive Criteria for *Bacillus anthracis* and *Brucella* spp.

Microorganism	MIC Interpretive Standard (µg/mL)		
	Susceptible (S)	Intermediate (I)	Resistant (R)
<i>Bacillus anthracis</i>	≤ 1	-	-
<i>Brucella</i> spp.			

Broth Microdilution performed in unsupplemented *Brucella* broth pH adjusted to 7.1 ± 0.1 for *Brucella* spp.⁵

For some organism/antimicrobial agent combinations, the absence or rare occurrence of resistant strains precludes defining results for categories other than "susceptible." For strains yielding results suggestive of a "nonsusceptible" category, organism identification and

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antimicrobial susceptibility test results should be confirmed.

Incubation in 5% CO₂ may be required for growth of some strains of *Brucella* spp. especially *B. abortus*.
 Incubation broth MIC tests in CO₂ may decrease the MIC of tetracyclines, usually by one doubling dilution.⁵

Microorganisms that are susceptible to tetracycline are generally susceptible to doxycycline.

Interpretive Criteria for *Bulkholderia mallei*, *Bulkholderia pseudomallei* and *Yersinia pestis*

Microorganism	MIC Interpretive Standard (µg/mL)		
	Susceptible (S)	Intermediate (I)	Resistant (R)
<i>Bulkholderia mallei</i>	≤ 4	8	≥ 16
<i>Bulkholderia pseudomallei</i>			
<i>Yersinia pestis</i>			

Microorganisms that are susceptible to tetracycline are generally susceptible to doxycycline.

Interpretive Criteria for *Franciscella tularensis*

Microorganism	MIC Interpretive Standard (µg/mL)		
	Susceptible (S)	Intermediate (I)	Resistant (R)
<i>Franciscella tularensis</i>	≤ 4	-	-

Broth Microdilutions performed in Cation-Adjusted Mueller-Hinton broth with 2% defined growth supplement for *Franciscella tularensis*⁵

For some organism/antimicrobial agent combinations, the absence or rare occurrence of resistant strains precludes defining results for categories other than "susceptible." For strains yielding results suggestive of a "nonsusceptible" category, organism identification and antimicrobial susceptibility test results should be confirmed.⁵

Microorganisms that are susceptible to tetracycline are generally susceptible to doxycycline.

A report of "Susceptible" indicates that the pathogen is likely to be inhibited if the antimicrobial compound in the blood reaches the concentrations usually achievable. A report of "Intermediate" indicates that the result should be considered equivocal, and, if the microorganism is not fully susceptible to alternative, clinically feasible drugs, the test should be repeated. This category implies possible clinical applicability in body sites where the drug is physiologically concentrated or in situations where high dosage of drug can be used. This category also provides a buffer zone which prevents small uncontrolled technical factors from causing major discrepancies in interpretation. A report of "Resistant" indicates that the pathogen is not likely to be inhibited if the antimicrobial compound in the blood reaches the concentrations usually achievable; other therapy should be selected.

Standardized susceptibility test procedures require the use of laboratory control microorganisms to control the technical aspects of the laboratory procedures. Standard tetracycline powder should provide the following MIC values:

Quality Control Ranges for MIC Broth Dilution Method

Microorganism	ATCC Number	Cation-Adjusted Mueller-Hinton (CAMHB) MIC (µg/mL)
<i>Escherichia coli</i>	25922	0.5 - 2
<i>Haemophilus influenzae</i>	49247	4 - 32*
<i>Neisseria gonorrhoeae</i>	49226	0.25 - 1†
<i>Pseudomonas aeruginosa</i>	27853	8 - 32
<i>Staphylococcus aureus</i>	29213	0.12 - 1
<i>Streptococcus pneumoniae</i>	49619	0.06 - 0.5‡

* Range applicable only to tests performed by broth microdilution method using *Haemophilus* Test Medium (HTM).^{1,3}

† Range applicable only to tests performed by agar dilution method using GC agar base with 1% defined growth supplement.^{1,3}

‡ Range applicable only to tests performed by broth microdilution method using Cation-Adjusted Mueller-Hinton broth with 2.5% to 5% lysed horse blood.^{1,3}

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Diffusion Techniques:

Quantitative methods that require measurement of zone diameters also provide reproducible estimates of the susceptibility of bacteria to antimicrobial compounds. One such standardized procedure^{2,3} requires the use of standardized inoculum concentrations. This procedure uses paper disks impregnated with 30-µg tetracycline or 30-µg doxycycline to test the susceptibility of microorganisms to doxycycline.

Reports from the laboratory providing results of the standard single-disk susceptibility test with a 30-µg tetracycline-class disk or the 30-µg doxycycline disk should be interpreted according to the following criteria:

Interpretive Zone Diameters for *Staphylococcus aureus*

Antimicrobial Agent	Zone Diameter (nearest whole mm)		
	Susceptible (S)	Intermediate (I)	Resistant (R)
Tetracycline	≥ 19	15 - 18	≤ 14
Doxycycline	≥ 16	13 - 15	≤ 12

Interpretive Zone Diameters for *Enterobacteriaceae*.

Antimicrobial Agent	Zone Diameter (nearest whole mm)		
	Susceptible (S)	Intermediate (I)	Resistant (R)
Tetracycline	≥ 15	12 - 14	≤ 11
Doxycycline	≥ 14	11 - 13	≤ 10
Minocycline	≥ 16	13 - 15	≤ 12

Interpretive Zone Diameters for *Acinetobacter* spp.

Antimicrobial Agent	Zone Diameter (nearest whole mm)		
	Susceptible (S)	Intermediate (I)	Resistant (R)
Tetracycline	≥ 15	12 - 14	≤ 11
Doxycycline	≥ 13	10 - 12	≤ 9
Minocycline	≥ 16	13 - 15	≤ 12

Interpretive Zone Diameters for *Haemophilus* spp.

Antimicrobial Agent	Zone Diameter (nearest whole mm)		
	Susceptible (S)	Intermediate (I)	Resistant (R)
Tetracycline	≥ 29	26 - 28	≤ 25

Interpretive criteria applicable only to tests performed by disk diffusion method using a 30-µg tetracycline-class disk and using *Haemophilus* Test Medium (HTM).^{2,3}

Microorganisms that are susceptible to tetracycline are generally susceptible to doxycycline.

Interpretive Zone Diameters for *Neisseria gonorrhoeae*

Antimicrobial Agent	Zone Diameter (nearest whole mm)		
	Susceptible (S)	Intermediate (I)	Resistant (R)
Tetracycline	≥ 38	31 - 37	≤ 30

Interpretive criteria applicable only to tests performed by disk diffusion method using a 30-µg tetracycline-class disk and using GC agar base with 1% defined growth supplement.^{2,3}
Zone diameters ≤ 19 mm may indicate a plasmid-mediated tetracycline-resistant *Neisseria gonorrhoeae* (TRNG) isolate. These TRNG strains should be confirmed by the dilution test (MIC ≥ 16 µg/mL).

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Microorganisms that are susceptible to tetracycline are generally susceptible to doxycycline.

Interpretive Zone Diameters for *Streptococcus pneumoniae*

Antimicrobial Agent	Zone Diameter (nearest whole mm)		
	Susceptible (S)	Intermediate (I)	Resistant (R)
Tetracycline	≥ 23	19 - 22	≤ 18

Interpretative criteria applicable only to tests performed by disk diffusion method using a 30-µg tetracycline-class disk and using Mueller-Hinton agar with 5% defibrinated sheep blood and incubated in 5% CO₂.^{2,3}

Microorganisms that are susceptible to tetracycline are generally susceptible to doxycycline.

Interpretation should be as stated above for results using dilution techniques. Interpretation involves correlation of the diameter obtained in the disk test with the MIC for tetracycline or doxycycline, respectively.

As with standardized dilution techniques, diffusion methods require the use of laboratory control microorganisms that are used to control the technical aspects of the laboratory procedures. For the diffusion technique, the 30-µg tetracycline-class disk or the 30-µg doxycycline disk should provide the following zone diameters in these laboratory test quality control strains:

Quality Control Zone Diameters for Disk Diffusion Method

Microorganism	ATCC Number	Zone Diameter (mm)		
		Tetracycline	Doxycycline	Minocycline
<i>Escherichia coli</i>	25922	18 - 25	18 - 24	19 - 25
<i>Haemophilus influenzae</i>	49247	14 - 22*	-	-
<i>Neisseria gonorrhoeae</i>	49226	30 - 42†	-	-
<i>Staphylococcus aureus</i>	25923	24 - 30	23 - 29	25 - 30
<i>Streptococcus pneumoniae</i>	49619	27 - 31‡	-	-

* Range applicable only to tests performed by disk diffusion method using a 30-µg tetracycline-class disk and using *Haemophilus* Test Medium (HTM).^{2,3}

† Range applicable only to tests performed by disk diffusion method using a 30-µg tetracycline-class disk and using GC agar base with 1% defined growth supplement.^{2,3}

‡ Range applicable only to tests performed by disk diffusion method using a 30-µg tetracycline-class disk and using Mueller-Hinton agar with 5% defibrinated sheep blood and incubated in 5% CO₂.^{2,3}

Anaerobic Techniques:

For anaerobic bacteria, the susceptibility to tetracycline as MIC's can be determined by standardized test methods.^{3,4} The MIC values obtained should be interpreted according to the following criteria:

Agar Dilution Interpretive Criteria for Anaerobes

Microorganism	MIC Interpretive Standard (µg/mL)		
	Susceptible (S)	Intermediate (I)	Resistant (R)
Anaerobes	≤ 4	8	≥ 16

Microorganisms that are susceptible to tetracycline are generally susceptible to doxycycline.

Interpretation is identical to that stated above for results using dilution techniques.

As with other susceptibility techniques, the use of laboratory control microorganisms is required to control the technical aspects of the laboratory standardized procedures. Standardized tetracycline powder should provide the following MIC values:

Quality Control Ranges for MIC Agar Dilution Method

Microorganism	ATCC Number	MIC (µg/mL)
<i>Bacteroides fragilis</i>	25285	0.125 - 0.5

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<i>Bacteroides thetaiotamicron</i>	29741	8 - 32
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Range applicable only to tests performed by the reference agar dilution method.

INDICATIONS AND USAGE

To reduce the development of drug-resistant bacteria and maintain effectiveness of Monodox and other antibacterial drugs, Monodox should be used only to treat or prevent infections that are proven or strongly suspected to be caused by susceptible bacteria. 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.

Doxycycline is indicated for the treatment of the following infections:

- Rocky mountain spotted fever, typhus fever and the typhus group, Q fever, rickettsialpox, and tick fevers caused by *Rickettsiae*.
- Respiratory tract infections caused by *Mycoplasma pneumoniae*.
- Lymphogranuloma venereum caused by *Chlamydia trachomatis*.
- Psittacosis (ornithosis) caused by *Chlamydia psittaci*.
- Trachoma caused by *Chlamydia trachomatis*, although the infectious agent is not always eliminated as judged by immunofluorescence.
- Inclusion conjunctivitis caused by *Chlamydia trachomatis*.
- Uncomplicated urethral, endocervical or rectal infections in adults caused by *Chlamydia trachomatis*.
- Nongonococcal urethritis caused by *Ureaplasma urealyticum*.
- Relapsing fever due to *Borrelia recurrentis*.

Doxycycline is also indicated for the treatment of infections caused by the following gram-negative microorganisms:

- Chancroid caused by *Haemophilus ducreyi*.
- Plague due to *Yersinia pestis* (formerly *Pasteurella pestis*).
- Tularemia due to *Francisella tularensis* (formerly *Pasteurella tularensis*).
- Cholera caused by *Vibrio cholerae* (formerly *Vibrio comma*).
- Campylobacter fetus infections caused by *Campylobacter fetus* (formerly *Vibrio fetus*).
- Brucellosis due to *Brucella species* (in conjunction with streptomycin).
- Bartonellosis due to *Bartonella bacilliformis*.
- Granuloma inguinale caused by *Calymmatobacterium granulomatis*.

Because many strains of the following groups of microorganisms have been shown to be resistant to doxycycline, culture and susceptibility testing are recommended.

Doxycycline is indicated for treatment of infections caused by the following gram-negative microorganisms, when bacteriologic testing indicates appropriate susceptibility to the drug:

- Escherichia coli*
- Enterobacter aerogenes* (formerly *Aerobacter aerogenes*)
- Shigella species*
- Acinetobacter species* (formerly *Mima species* and *Herellea species*)
- Respiratory tract infections caused by *Haemophilus influenzae*.
- Respiratory tract and urinary tract infections caused by *Klebsiella species*.

Doxycycline is indicated for treatment of infections caused by the following gram-positive microorganisms when bacteriologic testing indicates appropriate susceptibility to the drug:

- Upper respiratory infections caused by *Streptococcus pneumoniae* (formerly *Diplococcus pneumoniae*).
- Skin and skin structure infections caused by *Staphylococcus aureus*.

Anthrax due to *Bacillus anthracis*, including inhalational anthrax (post-exposure): to reduce the incidence or progression of disease following exposure to aerosolized *Bacillus anthracis*.

Doxycycline is not the drug of choice in the treatment of any type of staphylococcal infections.

When penicillin is contraindicated, doxycycline is an alternative drug in the treatment of the following infections:

- Uncomplicated gonorrhea caused by *Neisseria gonorrhoeae*.
- Syphilis caused by *Treponema pallidum*.
- Yaws caused by *Treponema pertenue*.
- Listeriosis due to *Listeria monocytogenes*.
- Vincent's infection caused by *Fusobacterium fusiforme*.
- Actinomycosis caused by *Actinomyces israelii*.
- Infections caused by *Clostridium species*.

In acute intestinal amebiasis, doxycycline may be a useful adjunct to amebicides.

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In severe acne, doxycycline may be useful adjunctive therapy.

CONTRAINDICATIONS

This drug is contraindicated in persons who have shown hypersensitivity to any of the tetracyclines.

WARNINGS

THE USE OF DRUGS OF THE TETRACYCLINE CLASS DURING TOOTH DEVELOPMENT (LAST HALF OF PREGNANCY, INFANCY, AND CHILDHOOD TO THE AGE OF 8 YEARS) MAY CAUSE PERMANENT DISCOLORATION OF THE TEETH (YELLOW-GRAY-BROWN). This adverse reaction is more common during long-term use of the drugs but has been observed following repeated short-term courses. Enamel hypoplasia has also been reported. TETRACYCLINE DRUGS, THEREFORE, SHOULD NOT BE USED IN THIS AGE GROUP, EXCEPT FOR ANTHRAX, INCLUDING INHALATIONAL ANTHRAX (POST-EXPOSURE), UNLESS OTHER DRUGS ARE NOT LIKELY TO BE EFFECTIVE OR ARE CONTRAINDICATED.

Clostridium difficile associated diarrhea (CDAD) has been reported with use of nearly all antibacterial agents, including Monodox, and may range in severity from mild diarrhea to fatal colitis. Treatment with antibacterial agents alters the normal flora of the colon leading to overgrowth of *C. difficile*.

C. difficile produces toxins A and B which contribute to the development of CDAD. Hypertoxin producing strains of *C. difficile* cause increased morbidity and mortality, as these infections can be refractory to antimicrobial therapy and may require colectomy. CDAD must be considered in all patients who present with diarrhea following antibiotic use. Careful medical history is necessary since CDAD has been reported to occur over two months after the administration of antibacterial agents.

If CDAD is suspected or confirmed, ongoing antibiotic use not directed against *C. difficile* may need to be discontinued. Appropriate fluid and electrolyte management, protein supplementation, antibiotic treatment of *C. difficile*, and surgical evaluation should be instituted as clinically indicated.

All tetracyclines form a stable calcium complex in any bone-forming tissue. A decrease in the fibula growth rate has been observed in prematures given oral tetracycline in doses of 25 mg/kg every six hours. This reaction was shown to be reversible when the drug was discontinued.

Results of animal studies indicate that tetracyclines cross the placenta, are found in fetal tissues, and can have toxic effects on the developing fetus (often related to retardation of skeletal development). Evidence of embryo toxicity has been noted in animals treated early in pregnancy. If any tetracycline is used during pregnancy or if the patient becomes pregnant while taking these drugs, the patient should be apprised of the potential hazard to the fetus.

The antianabolic action of the tetracyclines may cause an increase in BUN. Studies to date indicate that this does not occur with the use of doxycycline in patients with impaired renal function.

Photosensitivity manifested by an exaggerated sunburn reaction has been observed in some individuals taking tetracyclines. Patients apt to be exposed to direct sunlight or ultraviolet light should be advised that this reaction can occur with tetracycline drugs, and treatment should be discontinued at the first evidence of skin erythema.

PRECAUTIONS

General:

As with other antibiotic preparations, use of this drug may result in overgrowth of non-susceptible organisms, including fungi. If superinfection occurs, the antibiotic should be discontinued and appropriate therapy instituted.

Bulging fontanels in infants and benign intracranial hypertension in adults have been reported in individuals receiving tetracyclines. These conditions disappeared when the drug was discontinued.

Incision and drainage or other surgical procedures should be performed in conjunction with antibiotic therapy when indicated.

Prescribing Monodox in the absence of proven or strongly suspected bacterial infection or a prophylactic indication is unlikely to provide benefit to the patient and increases the risk of the development of drug-resistant bacteria.

Information for Patients:

All patients taking doxycycline should be advised:

- to avoid excessive sunlight or artificial ultraviolet light while receiving doxycycline and to discontinue therapy if phototoxicity (e.g., skin eruptions, etc.) occurs. Sunscreen or sunblock should be considered. (See WARNINGS.)
- to drink fluids liberally along with doxycycline to reduce the risk of esophageal irritation and ulceration. (See ADVERSE REACTIONS.)
- that the absorption of tetracyclines is reduced when taken with foods, especially those which contain calcium. However, the absorption of doxycycline is not markedly influenced by simultaneous ingestion of food or milk. (See Drug Interactions.)

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- that the absorption of tetracyclines is reduced when taking bismuth subsalicylate. (See Drug Interactions.)
- not to use outdated or poorly stored doxycycline.
- that the use of doxycycline might increase the incidence of vaginal candidiasis.

Diarrhea is a common problem caused by antibiotics which usually ends when the antibiotic is discontinued. Sometimes after starting treatment with antibiotics, patients can develop watery and bloody stools (with or without stomach cramps and fever) even as late as two or more months after having taken the last dose of the antibiotic. If this occurs, patients should contact their physician as soon as possible.

Patients should be counseled that antibacterial drugs including Monodox should only be used to treat bacterial infections. They do not treat viral infections (e.g., the common cold). When Monodox is prescribed to treat a bacterial 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 Monodox or other antibacterial drugs in the future.

Laboratory Tests:

In venereal disease when coexistent syphilis is suspected, a dark-field examination should be done before treatment is started and the blood serology repeated monthly for at least four months.

In long-term therapy, periodic laboratory evaluations of organ systems, including hematopoietic, renal, and hepatic studies should be performed.

Drug Interactions:

Because tetracyclines have been shown to depress plasma prothrombin activity, patients who are on anticoagulant therapy may require downward adjustment of their anticoagulant dosage.

Since bacteriostatic drugs may interfere with the bactericidal action of penicillin, it is advisable to avoid giving tetracyclines in conjunction with penicillin.

Absorption of tetracyclines is impaired by antacids containing aluminum, calcium, or magnesium, and iron-containing preparations.

Barbiturates, carbamazepine, and phenytoin decrease the half-life of doxycycline.

The concurrent use of tetracycline and methoxyflurane has been reported to result in fatal renal toxicity.

Concurrent use of tetracycline may render oral contraceptives less effective.

Drug/Laboratory Test Interactions:

False elevations of urinary catecholamine levels may occur due to interference with the fluorescence test.

Carcinogenesis, Mutagenesis, Impairment of Fertility:

Long-term studies in animals to evaluate the carcinogenic potential of doxycycline have not been conducted. However, there has been evidence of oncogenic activity in rats in studies with related antibiotics, oxytetracycline (adrenal and pituitary tumors) and minocycline (thyroid tumors). Likewise, although mutagenicity studies of doxycycline have not been conducted, positive results in *in vitro* mammalian cell assays have been reported for related antibiotics (tetracycline, oxytetracycline). Doxycycline administered orally at dosage levels as high as 250 mg/kg/day had no apparent effect on the fertility of female rats. Effect on male fertility has not been studied.

Pregnancy:

Teratogenic Effects.

Pregnancy Category D:

There are no adequate and well-controlled studies on the use of doxycycline in pregnant short-term, first trimester exposure. There are no human data available to assess the effects of long-term therapy of doxycycline in pregnant women such as that proposed for treatment of anthrax exposure. An expert review of published data on experiences with doxycycline use during pregnancy by TERIS - the Teratogen Information System - concluded that therapeutic doses during pregnancy are unlikely to pose a substantial teratogenic risk (the quantity and quality of data were assessed as limited to fair), but the data are insufficient to state that there is no risk.^a

A case-control study (18,515 mothers of infants with congenital anomalies and 32,804 mothers of infants with no congenital anomalies) shows a weak but marginally statistically significant association with total malformations and use of doxycycline anytime during pregnancy. (Sixty-three [0.19%] of the controls and 56 [0.30%] of the cases were treated with doxycycline.) This association was not seen when the analysis was confined to maternal treatment during the period of organogenesis (i.e., in the second and third months of gestation) with the exception of a marginal relationship with neural tube defect based on only two exposed cases.^b

A small prospective study of 81 pregnancies describes 43 pregnant women treated for 10 days with doxycycline during early first trimester. All mothers reported their exposed infants were normal at 1 year of age.^c

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Labor and Delivery:

The effect of tetracyclines on labor and delivery is unknown.

Nursing Mothers:

Tetracyclines are excreted in human milk, however, the extent of absorption of tetracyclines, including doxycycline, by the breastfed infant is not known. Short-term use by lactating women is not necessarily contraindicated; however, the effects of prolonged exposure to doxycycline in breast milk are unknown^d. Because of the potential for adverse reactions in nursing infants from doxycycline, a decision should be made whether to discontinue nursing or to discontinue the drug, taking into account the importance of the drug to the mother. (See WARNINGS.)

Pediatric Use:

See WARNINGS and DOSAGE AND ADMINISTRATION sections.

ADVERSE REACTIONS

Due to oral doxycycline's virtually complete absorption, side effects to the lower bowel, particularly diarrhea, have been infrequent. The following adverse reactions have been observed in patients receiving tetracyclines.

Gastrointestinal: Anorexia, nausea, vomiting, diarrhea, glossitis, dysphagia, enterocolitis, and inflammatory lesions (with monilial overgrowth) in the anogenital region. Hepatotoxicity has been reported. These reactions have been caused by both the oral and parenteral administration of tetracyclines. Rare instances of esophagitis and esophageal ulcerations have been reported in patients receiving capsule and tablet forms of drugs in the tetracycline class. Most of these patients took medications immediately before going to bed. (See DOSAGE AND ADMINISTRATION.)

Skin: Maculopapular and erythematous rashes. Stevens-Johnson syndrome, toxic epidermal necrolysis, and erythema multiforme have been reported. Exfoliative dermatitis has been reported but is uncommon. Photosensitivity is discussed above. (See WARNINGS.)

Renal Toxicity: Rise in BUN has been reported and is apparently dose related. (See WARNINGS.)

Hypersensitivity Reactions: Urticaria, angioneurotic edema, anaphylaxis, anaphylactoid purpura, serum sickness, pericarditis, and exacerbation of systemic lupus erythematosus.

Blood: Hemolytic anemia, thrombocytopenia, neutropenia, and eosinophilia have been reported with tetracyclines.

Other: Bulging fontanel in infants and intracranial hypertension in adults. (See PRECAUTIONS-General.)

When given over prolonged periods, tetracyclines have been reported to produce brown-black microscopic discoloration of the thyroid gland. No abnormalities of thyroid function are known to occur.

OVERDOSAGE

In case of overdosage, discontinue medication, treat symptomatically and institute supportive measures. Dialysis does not alter serum half-life, and it would not be of benefit in treating cases of overdosage.

DOSAGE AND ADMINISTRATION

THE USUAL DOSAGE AND FREQUENCY OF ADMINISTRATION OF DOXYCYCLINE DIFFERS FROM THAT OF THE OTHER TETRACYCLINES. EXCEEDING THE RECOMMENDED DOSAGE MAY RESULT IN AN INCREASED INCIDENCE OF SIDE EFFECTS.

Adults: The usual dose of oral doxycycline is 200 mg on the first day of treatment (administered 100 mg every 12 hours or 50 mg every 6 hours) followed by a maintenance dose of 100 mg/day. The maintenance dose may be administered as a single dose or as 50 mg every 12 hours. In the management of more severe infections (particularly chronic infections of the urinary tract), 100 mg every 12 hours is recommended.

For pediatric patients above eight years of age: The recommended dosage schedule for pediatric patients weighing 100 pounds or less is 2 mg/lb of body weight divided into two doses on the first day of treatment, followed by 1 mg/lb of body weight given as a single daily dose or divided into two doses, on subsequent days. For more severe infections up to 2 mg/lb of body weight may be used. For pediatric patients over 100 pounds the usual adult dose should be used.

Uncomplicated gonococcal infections in adults (except anorectal infections in men): 100 mg, by mouth, twice a day for 7 days. As an alternate single visit dose, administer 300 mg stat followed in one hour by a second 300 mg dose.

Acute epididymo-orchitis caused by *N. gonorrhoeae*: 100 mg, by mouth, twice a day for at least 10 days.

Primary and secondary syphilis: 300 mg a day in divided doses for at least 10 days.

Uncomplicated urethral, endocervical, or rectal infection in adults caused by *Chlamydia trachomatis*: 100 mg, by mouth, twice a day for

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at least 7 days.

Nongonococcal urethritis caused by *C. trachomatis* and *U. urealyticum*: 100 mg, by mouth, twice a day for at least 7 days.

Acute epididymo-orchitis caused by *C. trachomatis*: 100 mg, by mouth, twice a day for at least 10 days.

Inhalational anthrax (post-exposure): ADULTS: 100 mg of doxycycline, by mouth, twice a day for 60 days. CHILDREN: weighing less than 100 pounds (45 kg); 1 mg/lb (2.2 mg/kg) of body weight, by mouth, twice a day for 60 days. Children weighing 100 pounds or more should receive the adult dose.

When used in streptococcal infections, therapy should be continued for 10 days.

Administration of adequate amounts of fluid along with capsule and tablet forms of drugs in the tetracycline class is recommended to wash down the drugs and reduce the risk of esophageal irritation and ulceration. (See ADVERSE REACTIONS.) If gastric irritation occurs, doxycycline may be given with food. Ingestion of a high fat meal has been shown to delay the time to peak plasma concentrations by an average of one hour and 20 minutes. However, in the same study, food enhanced the average peak concentration by 7.5% and the area under the curve by 5.7%.

HOW SUPPLIED

MONODOX® 50 mg Capsules have a white opaque body with a yellow opaque cap. The capsule bears the inscription "MONODOX 50" in brown and "M 260" in brown. Each capsule contains doxycycline monohydrate equivalent to 50 mg doxycycline.

MONODOX® 50 mg is available in:

Bottles of 100 capsulesNDC 16110-260-06

MONODOX® 75 mg Capsules have a white opaque body with a brown opaque cap. The capsule bears the inscription "MONODOX 75" in black and "M 075" in black. Each capsule contains doxycycline monohydrate equivalent to 75 mg doxycycline.

MONODOX® 75 mg is available in:

Bottles of 100 capsulesNDC 16110-075-01

MONODOX® 100 mg Capsules have a yellow opaque body with a brown opaque cap. The capsule bears the inscription "MONODOX 100" in white and "M 259" in brown. Each capsule contains doxycycline monohydrate equivalent to 100 mg of doxycycline.

MONODOX® 100 mg is available in:

Bottles of 50 capsulesNDC 16110-259-04

Bottles of 250 capsulesNDC 16110-259-07

STORE AT 20° - 25°C (68° - 77°F). [SEE USP CONTROLLED ROOM TEMPERATURE.]

DISPENSE IN A TIGHT LIGHT-RESISTANT CONTAINER AS DEFINED IN THE USP/NF.

ANIMAL PHARMACOLOGY AND ANIMAL TOXICOLOGY

Hyperpigmentation of the thyroid has been produced by members of the tetracycline class in the following species: in rats by oxytetracycline, doxycycline, tetracycline PO₄, and methacycline; in minipigs by doxycycline, minocycline, tetracycline PO₄, and methacycline; in dogs by doxycycline and minocycline; in monkeys by minocycline.

Minocycline, tetracycline PO₄, methacycline, doxycycline, tetracycline base, oxytetracycline HCl and tetracycline HCl were goitrogenic in rats fed a low iodine diet. This goitrogenic effect was accompanied by high radioactive iodine uptake. Administration of minocycline also produced a large goiter with high radioiodine uptake in rats fed a relatively high iodine diet.

Treatment of various animal species with this class of drugs has also resulted in the induction of thyroid hyperplasia in the following: in rats and dogs (minocycline), in chickens (chlortetracycline) and in rats and mice (oxytetracycline). Adrenal gland hyperplasia has been observed in goats and rats treated with oxytetracycline.

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