INTERIM FORMULARY UPDATE Draft

The following recommendations, made at the February 10, 2006 meeting of the Executive Formulary Committee, are approved:

Product(s) **approved to be added** to the DADS/DSHS Drug Formulary based on the Formulary Review:

Generic Name	Brand Name	Dosage Form	Classification	
Valsartan	Diovan®	Tablet: 40 mg, 80 mg,	Angiotensin II Receptor	
		160 mg, 320 mg	Blocking Agent (ARBs)	
Olmesartan	Benicar®	Tablet: 5 mg, 20 mg,	Angiotensin II Receptor	
		40 mg	Blocking Agent (ARBs)	

Product(s) **approved to be added** to the DADS/DSHS Drug Formulary based on the Sectional Reviews for Gastrointestinal, Genitourinary Agents: **Not completed will be presented at next meeting.**

Dosage strength/formulations **recommended to be deleted** from the DADS/DSHS Drug Formulary based on the Sectional Review: **Not completed will be presented at next meeting.**

Previously, the Executive Formulary Committee requested input from the field regarding the following drugs proposed for deletion from the DADS/DSHS Drug Formulary. Based on the field's response, the following drugs are deleted from the Drug Formulary.

Products **deleted** from the DADS/DSHS Drug Formulary based on these previous Sectional Reviews: for Infectious Disease Agents:

Generic Name	Brand Name	Dosage Form	Dosage Forms
			Still Available
Cefoperazone	Cefobid®	Infusion, premixed in dextrose: 1 g, 2 g	None
Chloroquine	Aralen®	Tablet: 250 mg, 500 mg	None
Cloxacillin	Cloxapen®,	Capsule: 250 mg, 500 mg	None
	Tegopen®	Powder for oral suspension: 125 mg/5 ml	
Ethionamide		Tablet, sugar-coated: 250 mg	None
Pentamidine	Pentam®	Inhalation:300 mg	None
		Powder for injection: 300 mg	
Pyrantel	Antiminth®	Capsule: 180 mg	None
		Liquid, oral: 50 mg/ml	
		Suspension, oral: 50 mg/ml	
Thiabendazole	Mintezol®	Suspension, oral: 500 mg/5 ml	None
		Tablet, chewable: 500 mg	
Ticarcillin	Ticar®	Powder for injection: 1 g, 3 g, 6 g, 20 g, 30 g	None
Ticarcillin/	Timentin®	Powder for injection: 3.1 g	None
clavulanate			

Other recommendations:

- All drug audit criteria be changed from hepatic function panel to hepatic function testing and TDMHMR Drug Formulary to DSHS/DADS Drug Formulary.
- The statement "Monitor for emergence of suicidal ideation or behavior" be added to all antidepressant audit criteria as a monitoring parameter.

Other **recommendation(s)/addition(s)/revisions(s)** to the DADS/DSHS Drug Formulary:

- Add ARBs valsartan and olmesartan to the reserve category. Criteria for use of the ARBs will be: Prior failure to ACE inhibitor therapy due to intolerable side effects.
- Remove divalproex ER from the reserve category.

Approved:

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Stever P. Shon, MD

Date: April 10, 2006