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Implementation Of A Guideline-Based Nontuberculous Mycobacteria Management Algorithm

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Background

- Nontuberculous mycobacterial (NTM) disease is a challenge to manage in patients with cystic fibrosis (CF).
- Diagnosis of NTM pulmonary disease is complex.
- Effective treatment requires long term, multi-drug therapy delivered by several routes.
- Consensus recommendations published in 2016 were developed to guide CF providers in NTM screening, diagnosis and management.

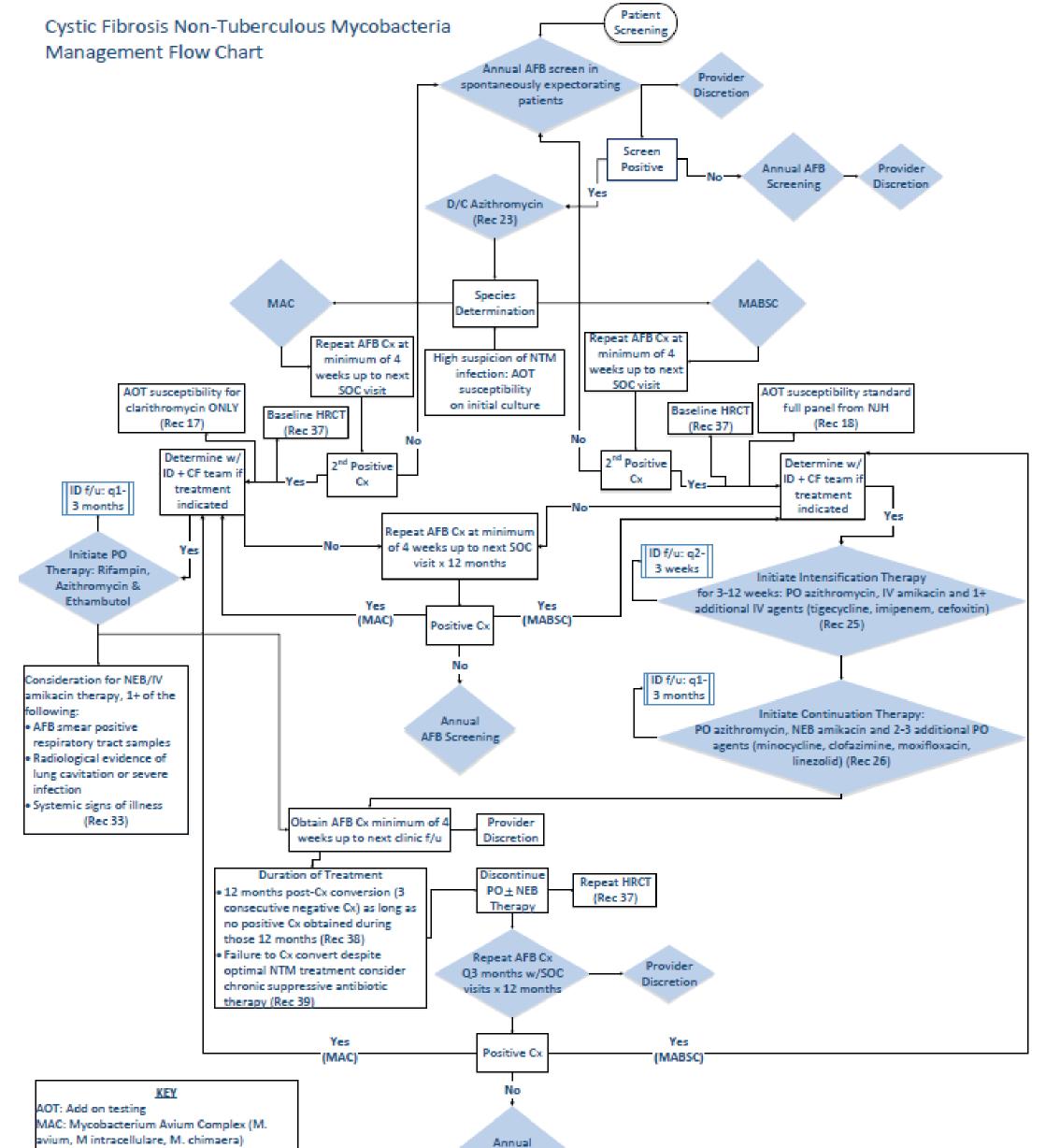
Primary Outcome

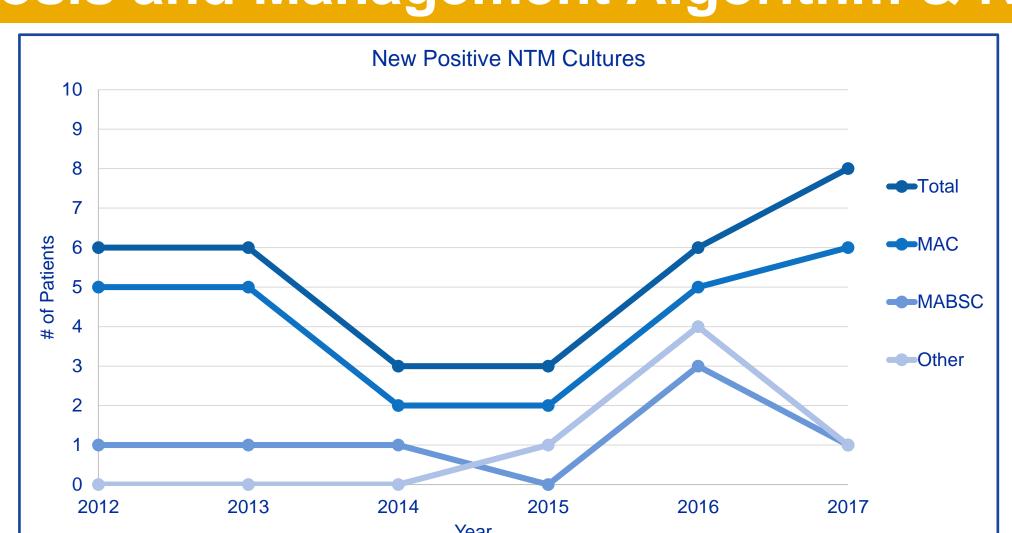
Children's Mercy – Kansas City (CMKC) developed an NTM working group to facilitate implementation of standardized NTM management.

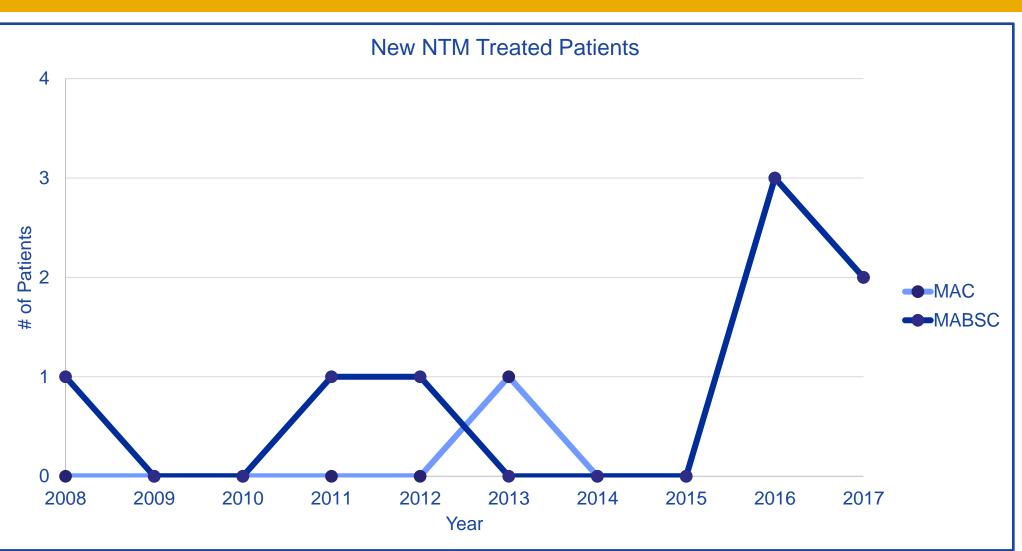
Methods

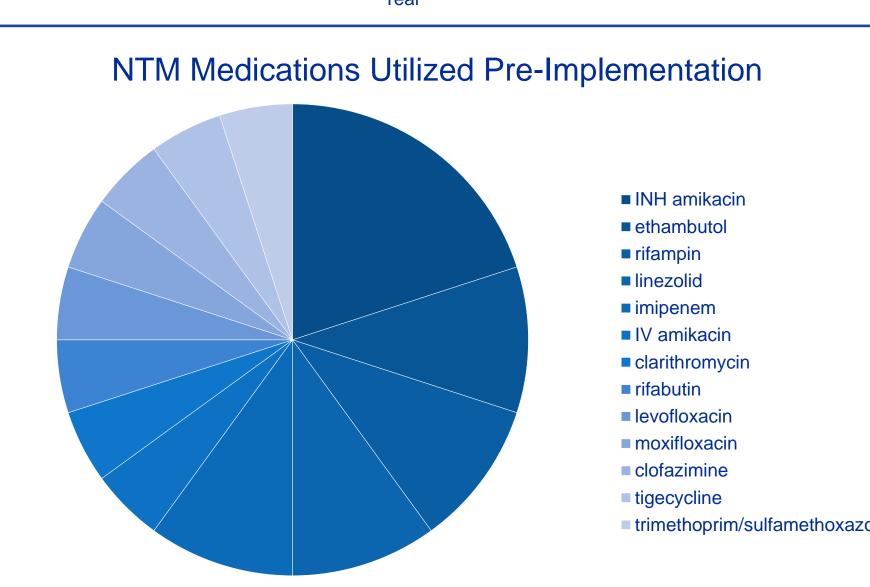
- NTM working group: Pulmonologist (CF Center Director), Infectious Diseases specialist, nurse practitioner (CF Center Coordinator), and pharmacist.
- Sought expertise from the Director of Microbiology Laboratory regarding susceptibility testing.
- Developed NTM Management Guide from guidelines and other available literature.
 - Diagnosis and management algorithm
 - Medication resource table

Diagnosis and Management Algorithm & Results



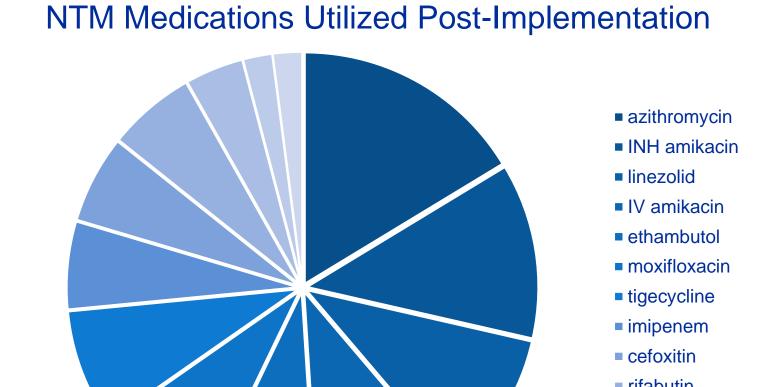








- Laboratory abnormalities
 Bronchospasm
- GI intolerance
- Musculoskeletal
- Peripheral neuropathyOtotoxicity
- abnormalitiesPhotosensitivity



Post-Implementation Identified ADRs

rifampin

minocycline

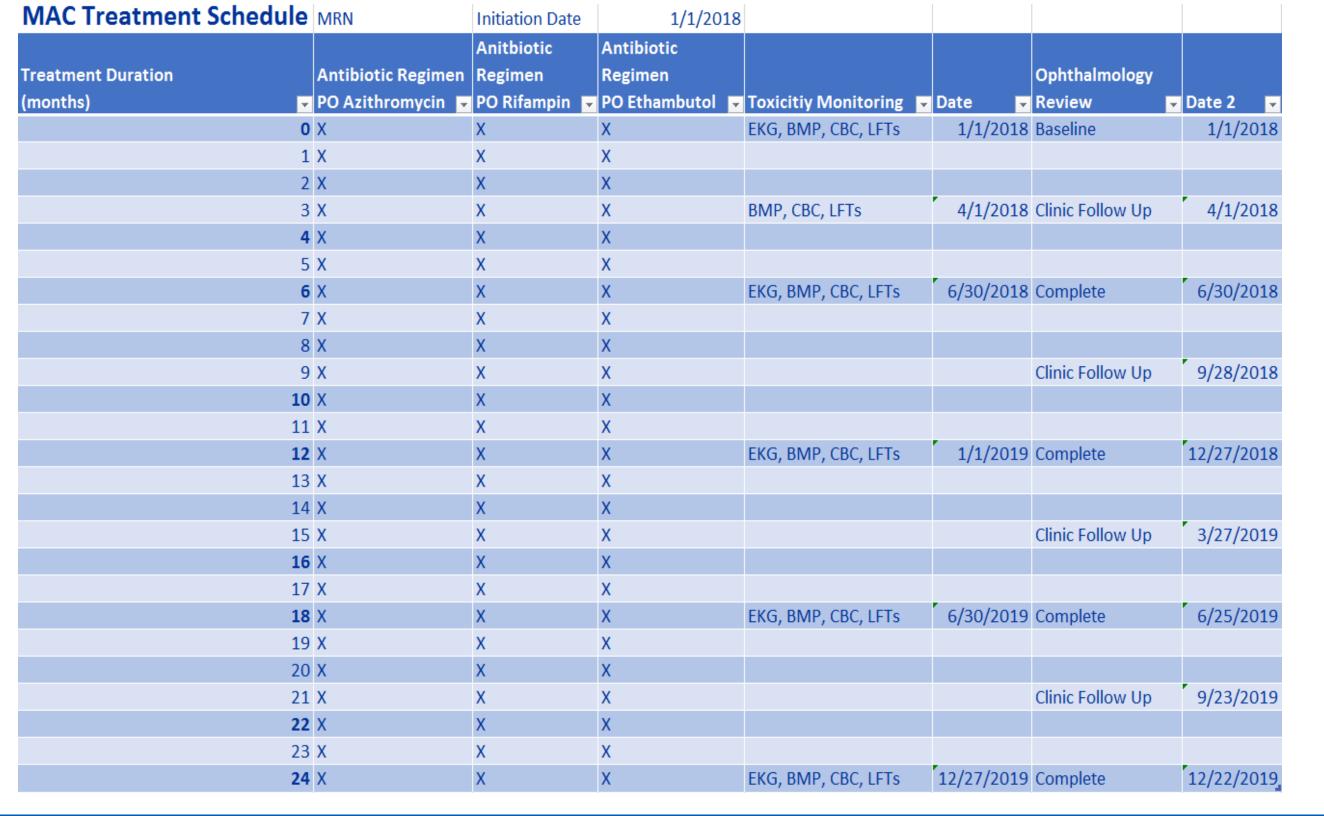
- Laboratory abnormalities
- Gl intolerance
- Ototoxicity

Medication Resource Table

RCT: High-resolution CT

UH: National Jewish Health

Agent	Indication	Dosing	Monitoring
			Serum amikacin levels, BMPs: weekly
			Audiograms:
Secoli a alte		NEB: 250 BID or 500 mg daily	IV: baseline, then monthly x 1 month post-IV discontinuation
Amikacin		IV: 15-30 mg/kg/dose once daily; max dose: 1500 mg	NEB: baseline, then every 6 months
-141	MAC	10.10	EKG: baseline, then every 6 months
Azithromycin		10-12 mg/kg/dose PO once daily; max dose: 500 mg	LFTs: baseline, every 6 months
Cefoxitin	MABSC	50 mg/kg/dose IV q 8 hours; max dose: 4000 mg/dose	CBC, BMP, LFTs: weekly
	MAC		Skin discoloration
Clofazimine	MABSC	1-2 mg/kg/dose PO once daily; max dose: 100 mg	CBC, BMP, LFTs: baseline, then monthly
			Ophthalmic exam: baseline, then monthly up to next clinic follow up; every 6
	MAC	Initial: 15 mg/kg/dose PO once daily; max dose: 1600 mg	months complete
thambutol		Re-treatment (max): 25 mg/kg/dose	BMP, CBC, LFTs: baseline, then every 3 months x 2, then every 6 months
mipenem/	MABSC		
Cilastatin	IVIADOC	100 mg/kg/day IV every 6 hours; max dose: 1000 mg	CBC, BMP, LFTs: weekly
			CBC: baseline, weekly on IV therapy then every 2 weeks
	MABSC	< 12 years: 10 mg/kg/dose PO/IV q8 hours	BMP, LFTs: baseline, weekly on IV therapy then monthly
inezolid		>12 years: 10 mg/kg/dose PO/IV q 12 hours; max dose: 600 mg	Ophthalmic exam: baseline, every 6 months
/linocycline	MABSC	2 mg/kg/dose PO twice daily; max dose: 100 mg	CBC, BMP, LFTs: baseline, then every 3 months x 2, then every 6 months
	MADCC		EKG: baseline, then every 6 months
Moxifloxacin	MABSC	7.5-10 mg/kg/dose PO once daily; max dose: 400 mg	CBC, HFP: baseline, then every 3 months x 2, then every 6 months
Rifampin	MAC	10-20 mg/kg/dose PO once daily; max dose: 600 mg	CBC, LFTs: baseline, then every 3 months x 2, then every 6 months
Sulfamethoxazole		TMP: 15-20 mg/kg/dose PO/IV q 6-12 hours	
rimethoprim	MABSC	max: TMP 1920 mg/day	CBC, BMP, LFTs: baseline, then every 3 months x 2, then every 6 months
		8-11 years: 1.2 mg/kg/dose IV q 12 hours; max dose: 50 mg	
igecycline	IIVIABSU		CBC, BMP, LFTs: weekly



Conclusion

- Post-implementation results demonstrate improved consistency in selection of medications and fewer adverse drug reactions despite treating more patients.
- The Medication Resource Table provides consistency in monitoring among pulmonary providers and helps families and ancillary disciplines plan monitoring visits and coordination of care.
- The NTM Management Guide allows implementation of NTM guidelines into clinical practice, standardization of our NTM management and improvement of patient care.

The authors of have no relevant disclosures.







