

Version 10.			IIVIA			MEDIC.	BIA UNIVERSITY AL CENTER
(mmb2225@c approving the	cumc.co study b	lumbia.edu) in	the O _l	phthalmo ees. Pleas	se include a copy of	or to th	autista e Clinical Trials Office dy Protocol, Case Report
					ualtrics.com/jfe/forr ESEARCH COORDINA		pAL2qDRniIk5Bs
		Name			Email		Phone Number
Principal Invest	tigator						
Study Coordina	ator						
Administrator							
Study Title:							
IRB #:		Estir	nated	number	of patients to be enro	olled in s	study:
Type of Study:	NIH Fun	nded 🔲 Indust	ry Tria	al 🗌 Sp	oonsor		
Study Start Dat	e:			Stud	y End Date		
Will you need	certifica	ation for refract	ion (v	isual acui	ty), ocular testing (vi	sual fiel	d), imaging (OCT)? Yes
Will this study	utilize a	Reading Center	? Yes	☐ No	Name of Rea	ading Ce	No enter:
Please identify	procedu	res that will us	e a Re	ading Ce	nter below.		
Requested Exam				Patient for Duration of Study		Needs copies, backup, reading center transmission for images or special requests?	
TO BE COMPLETE	D BY OPI	HTHALMOLOGY FO	R FEES				
CPT Code Procedure (both eyes) Price				<u> </u>	Additional Comments:		
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Start-up Fee

Administrative

^{*}There will be a start-up fee in the amount of \$360 to cover up to 4 hours of time for the clinical research coordinator to complete the cost estimate form; assist to schedule study subjects for eye exams, testing, and imaging; and answer email queries if needed. Any additional services required to conduct the study above 4 hours will be invoiced at \$90 per hour.

^{**}Excluding NIH funded clinical trials.