

Division of Human Genetics Level 3, Wernher and Beit North Institute of Infectious Disease and Molecular Medicine Faculty of Health Sciences University of Cape Town Observatory, 7925 South Africa

## **CONFIRMATION OF DIAGNOSIS**To be completed by the eye specialist – PLEASE PRINT CLEARLY

Name of	Patient									
Date of Birth: YYYY-MM-DD Tel/Cel					ell	Email				
Address:										
Gender:	М	F	Ethnic Gr	oup	Asian	Black	Coloured	Indian	White	
In my opinion the patient has one of the following conditions:										
RETINITIS PIGMENTOSA					Diffus	e Form				
						Sectoral (regional) form				
	USHER SYNDROME					Type I – profound deafness				
	(RP & congenital hearing loss)					Type II – severe deafness				
	MACULAR DEGENERATION					Age-related MD - Wet				
						- Dry				
					Best I	Disease	<b>,</b>			
					Cone	Cone & Rod Dystrophy				
						Sorsby Fundus Dystrophy				
						Pattern Dystrophy				
						Stargardt Disease				
						Fundus Flavimaculatus				
Other retinal disorder (specify):  MODE OF INHERITANCE  Dominant Recessive X-Linked Isolated Case Unknown but familial  Age of Onset: years  Progression of disease:  Other clinical features:										
Tooto				ıal	Kino	tic Visual				
performed:		ERG/E	OG	Acui			Fields	ОСТ		
Fluorescein Angiogram Colour Fundus Photographs										
Other family members affected:										
Name of Doctor:						Sig	nature:			
Date:	YYYY	- M M - D	D Tel.	. ( )			Email			

PLEASE RETURN THE COMPLETED FORM TO: retina@uct.ac.za

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