

## PATIENT ELIGIBILITY ASSESSMENT SHEET- FOR CLINICIANS

## Familial Pituitary Tumour Syndromes: identification of AIP, MEN-1, p27 mutations/deletions and other novel genes - An Australasian Perspective

	Patient initials: Patient D.O.B.	
	Doctor's name: Practice Address:	
	Phone: ( )  Fax: ( )	
	Email:	
ELI	LIGIBILITY CRITERIA (Please circle appropriate criteria)	
A)	) MEN-1 gene mutation positive from previous testing (with or without a pituitary tumour)  OR	
B)	Pituitary tumour diagnosed < 40 years of age  OR	
C)	<ul> <li>Pituitary tumour diagnosed &gt; 40 years of age <u>PLUS</u> family history of pituitary tumour (at ar OR</li> </ul>	ıy age)
D)	<ul> <li>Pituitary tumour diagnosed &gt; 40 years of age <u>PLUS</u> patient has other MEN-1 related neop OR</li> </ul>	lasia(s)
E)	) Pituitary tumour occurring in a patient > 40 years <u>PLUS</u> family member with other MEN-1 r	elated neoplasia(s)§
For	r patients in categories B to E please indicate whether:	
□ <b>N</b>	MEN-1 gene mutation negative (genetic testing previously undertaken)	
□ <b>N</b>	No genetic testing undergone as yet	
§ <b>T</b> y	Typical MEN1 related neoplasias: Primary hyperparathyroidism, anterior pituitary adenoma umours (e.g., gastrinoma, insulinoma), adrenal adenoma, bronchial carcinoid, thymic carcinoid	, GEP d.
EXC	(CLUSION CRITERIA:	
	any of the following apply:	

## **EXCLUSION C**

- ☐ Age less than 18 years
- ☐ Unable to give independent informed consent
- ☐ Unable to understand and/or read English
- ☐ Serious medical or psychiatric illness due to which the study might place undue burden on the participant or prevent them from completing the study

Please return this form with your patient's signed consent form via email or fax, to:

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