

VON HIPPEL LINDAU SYNDROME SCREENING PROTOCOL

Affected Individuals	
Physical examination & history	Annually
Ophthalmoscopy (direct & indirect), fluorescein angiography or angiography	Annually
Cranial + spinal MRI	Every 3 years from 18 until 60
Abdominal ultrasound	Annually - not fasting
Abdominal CT	Every 3 years – not fasting
Endocrine screen including <ul style="list-style-type: none"> • Metanephrines + normetanephrines • pancreatic peptides (fasting) - • glucose • insulin • C peptide 	Annually
Blood Pressure (erect and supine)	

Individuals at 50% risk	
Physical examination & history	Annually, from age 10
Ophthalmoscopy (direct & indirect), fluorescein angiography or angiography	Annually from age 10-60
Cranial + spinal MRI	Every 3 years from 18 until 40, then every 5 years until 60
Abdominal ultrasound	Annually from age 20-65
Abdominal CT	Every 3 years from age 20-65
Endocrine screen including <ul style="list-style-type: none"> • Metanephrines + normetanephrines • pancreatic peptides (fasting) - • glucose • insulin • C peptide 	Annually, from age 10
Blood Pressure (erect and supine)	

Protocol prepared by: South East of Scotland Clinical Genetic Service, 2001