Strong	Not Strong/Some	No Relevant Family History
Personal or family (1st or 2nd degree) history of Breast cancer ≤45 Breast cancer 45-50 w/ another breast cancer Triple neg breast cancer ≤60 Ovarian cancer Pancreatic cancer Metastatic prostate cancer Colon cancer <50 Other GI cancer <50 Uterine cancer <50 Brain cancer <20 Kidney <50 Eye cancer	Presence of these cancers (in 1 st or 2 nd), not strong Breast cancer Uterine cancer Colon cancer Non-metastatic Prostate cancer Sarcoma Other GI cancer Kidney Brain Leukemia	Lack of "strong" or "not strong" criteria Lymphoma/myeloma Liver cancer Bladder cancer Thyroid cancer Lung cancer Oral cancer Cervical cancer Testicular cancer Skin cancer (basal or squamous included)
High suspicion of cancer syndrome -3+ Lynch-related on same side of family -Breast cancer 46-50 with another breast at any age -2+ kidney 50+ on same side -sarcoma <45 plus other Call patient if personal history of: Breast cancer <45 Ovarian cancer Breast and ovarian cancer Colon cancer <49 Uterine cancer <49 Known familial mutation	Melanoma 3 rd who would meet strong	

Information is Power Cancer Syndrome Appendix

High Cancer Syndrome Suspicion if they mention:	Lynch syndrome (NCCN)	
Adrenocortical tumor	Cancers: colon, endometrial, ovarian, stomach, sebaceous	
Rhabdomyoma	adenoma, urinary tract, small bowel, brain, pancreas,	
Male breast cancer	prostate, hepatobiliary	
Pheochromocytoma/paraganglioma	Personal history 2+ Lynch cancers Personal halon (and assertial + 18 on 28 relative with 16).	
Hemangioblastoma	 Personal colon/endometrial + 1° or 2° relative with LS- related <50 	
Sertoli cell tumor	Telated <50	
Wilms tumor		
• 10+ polyps	1.504/6	
HBOC (NCCN)	MEN1 (GeneReviews)	
Cancers: breast, ovarian, prostate, pancreatic2 primary breast cancers	 Tumors: parathyroid, pituitary, gastro-entero-pancreatic tract 	
 More than one 1° or 2° relative with breast cancer + breast cancer <50 or other HBOC cancer 	At least one 1° relative with at least one of the tumors	
MEN2 (GeneReviews)	Gorlin syndrome (ACMG)	
Tumors: Medullary thyroid carcinoma, pheochromocytoma,	• 5+ BCC <30	
parathryroid adenoma	 BCC <30 + 1° relative with 5+ BCC <30 	
 2+ tumors in single person or 1° and 2° relatives 		
Cowden/PTEN (NCCN)	FAMMM (ACMG)	
Cancers: breast, endometrial, thyroid, colon, renal cell	3+ relatives with melanoma and/or pancreatic cancer	
carcinoma	3+ primary melanoma in same person	
 Relative with diagnosis + breast, endometrial, thyroid 	Melanoma and pancreatic cancer in one person	
Familial Adenomatous Polyposis (NCCN)	Xeroderma Pigmentosa (GeneReviews)	
Desmoid tumor, hepatoblastoma, cribiriform thyroid cancer	• Skin cancer <10	
Li-Fraumeni (NCCN)	BAP1 (GeneReviews)	
 Cancers: breast, brain, sarcoma, osteosarcoma, adrenocortical carcinoma, leukemia 	 Cancers: uveal melanoma, mesothelioma, melanoma, renal cell carcinoma, BCC 	
• 1° relative cancer <45y + 1° or 2° with cancer <45 or sarcoma	• 2+ BAP1 cancers	
	 Personal history + 1 BAP1 tumor in 1° or 2° relative 	
Peutz-Jeghers	Hereditary leiomyomatosis and renal cell cancer (ACMG)	
Cancers: breast, colon, stomach, pancreatic, lung, ovarian and	Leiomyoma + renal cell carcinoma	
tests (sex cord/Sertoli cell tumors), testes		
Criteria: 2+ PJS hamartomatous GI polyps + family history		