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**National Cancer Institute**  
at the National Institutes of Health

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### NCI Cancer Genetics Services Directory: Application Form

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**1. Please provide your full name, including middle initial, and the complete address, telephone number, fax number, and email address where you can be contacted for annual verification of your information:**

\* Name:

\* Institution:

\* Address:

\* Telephone Number:

Fax Number:

\* Email Address:

\* Publish your email address in the NCI Cancer Genetics Services Directory?  
Yes  No

Web site URL:

Do you provide services at this location?  
Yes  No

Web site URL:

Do you provide services at this location?  
Yes  No

If no, please provide information for service location:

\* Institution:

\* Service Address:

\* Telephone Number:

If you want additional locations, please enter them below (maximum of four):

**2. \* What type of health care professional are you?**

Physician (M.D., D.O., or foreign equivalent)  
 Geneticist (Ph.D.)  
 Genetic Counselor (M.S., M.Sc., M.A., C.G.C.)  
 Nurse (R.N., B.S.N., M.S.N., M.S., M.A., Ph.D.)  
 Clinical Social Worker (M.S.W., D.S.W.)  
 Clinical Psychologist (Ph.D., Psy.D.)  
 Other

Please specify degree(s):

Provide professional license and/or national certification number and state:

**3a. \* What is/are your specialties? (Choose one or more)**

Clinical Biochemical Genetics  
 Clinical Cytogenetics  
 Clinical Genetics  
 Clinical Molecular Genetics  
 Genetic Counseling  
 Gynecologic Oncology  
 Hematology  
 Medical Biochemical Genetics

Clinical Genetics  
 Clinical Molecular Genetics  
 Genetic Counseling  
 Gynecologic Oncology  
 Hematology  
 Medical Biochemical Genetics  
 Medical Genetics  
 Medical Oncology  
 Molecular Genetic Pathology  
 Oncology Nursing  
 Pediatric Hematology-Oncology  
 Radiation Oncology

**3b. \* Are you board certified in your specialty?**  
 Yes  No   
 If yes, please specify specialty board:   
 If no, are you board eligible? Yes  No   
 If yes, please provide the year you became eligible:

**4. What specific training or professional experience do you have in cancer genetics?**  
**Please include information about all of the following that apply:**

Citations for relevant publications:

Examples of relevant continuing education or graduate courses:

Clinical preceptorships taken:

Investigator (or other research professional) on genetics clinical trials:

Number of patients counseled per year about genetic susceptibility to cancer:   
 Number of years of experience:

**5. \* Are you a member of or affiliated with an interdisciplinary team with substantial expertise in cancer genetics?**  
 Yes  No

**6. \* For which of the following do you or members of your team provide expertise in relation to cancer genetics? (Choose one or more)**

**expertise in cancer genetics?**  
 Yes  No

**6. \* For which of the following do you or members of your team provide expertise in relation to cancer genetics? (Choose one or more)**

Patient genetics education  
 Patient cancer risk assessment  
 Appropriate pre- and post-test counseling and informed consent (including ethical, legal, social issues related to testing and disclosure of test results)  
 Genetic susceptibility testing (including information on limitations, specific tests available, and regulations concerning testing procedures such as CLIA and CAP/ACMG)  
 Follow-up plan of care (including medical care, psychological support, and counseling about options for prevention or early detection guidelines)

**7. \* Do you currently provide professional services to individuals or families seeking familial cancer risk counseling or genetic susceptibility testing? (Please note that if you do not provide services, you cannot be added to the directory.)**  
 Yes  No

**8. \* Are you willing to accept calls or email from individuals seeking familial cancer risk counseling and/or genetic susceptibility testing?**  
 Yes  No

**9. \* Are there restrictions or limitations to services provided (i.e., a person must be eligible for a clinical trial in order to receive services)?**  
 Yes, Explain:   
 No

**10. \* Please verify the familial cancer predisposing syndromes for which you provide services. A list of cancer sites and types associated with each syndrome will also be provided for searching in the directory. (Choose one or more)**

<input type="checkbox"/> Ataxia-telangiectasia	<input type="checkbox"/> Multiple endocrine neoplasia type 1
<input type="checkbox"/> Basal cell nevus syndrome	<input type="checkbox"/> Multiple endocrine neoplasia type 2
<input type="checkbox"/> Beckwith-Wiedemann syndrome	<input type="checkbox"/> Multiple myeloma, familial
<input type="checkbox"/> Birt-Hogg-Dubé syndrome	<input type="checkbox"/> Neuroblastoma, hereditary
<input type="checkbox"/> Bloom syndrome	<input type="checkbox"/> Neurofibromatosis type 1
<input type="checkbox"/> Breast/ovarian cancer, hereditary (BRCA1)	<input type="checkbox"/> Neurofibromatosis type 2
<input type="checkbox"/> Breast/ovarian cancer, hereditary (BRCA2)	<input type="checkbox"/> Nijmegen breakage syndrome
<input type="checkbox"/> Carney complex, types I and II	<input type="checkbox"/> Pancreatic cancer, hereditary
<input type="checkbox"/> Chordoma, familial	<input type="checkbox"/> Paraganglioma, hereditary
<input type="checkbox"/> Colon cancer, hereditary non-polyposis - Lynch syndrome	<input type="checkbox"/> Peutz-Jeghers syndrome
<input type="checkbox"/> Costello syndrome	<input type="checkbox"/> Polyposis, familial adenomatous
<input type="checkbox"/> Cowden syndrome	<input type="checkbox"/> Polyposis, familial juvenile
<input type="checkbox"/> Dyskeratosis congenita	<input type="checkbox"/> Polyposis, MYH-associated
<input type="checkbox"/> Esophageal cancer with tylosis	<input type="checkbox"/> Prostate cancer, hereditary
	<input type="checkbox"/> Renal cell cancer, hereditary

- |  |   |
|--|---|
| <input type="checkbox"/> Choroideroma, familial                                  | <input type="checkbox"/> Paraganglioma, hereditary                |
| <input type="checkbox"/> Colon cancer, hereditary non-polyposis - Lynch syndrome | <input type="checkbox"/> Peutz-Jeghers syndrome                   |
| <input type="checkbox"/> Costello syndrome                                       | <input type="checkbox"/> Polyposis, familial adenomatous          |
| <input type="checkbox"/> Cowden syndrome   | <input type="checkbox"/> Polyposis, familial juvenile             |
| <input type="checkbox"/> Dyskeratosis congenita                                  | <input type="checkbox"/> Polyposis, MYH-associated                |
| <input type="checkbox"/> Esophageal cancer with tylosis                          | <input type="checkbox"/> Prostate cancer, hereditary              |
| <input type="checkbox"/> Exostosis, hereditary multiple                          | <input type="checkbox"/> Renal cell cancer, hereditary            |
| <input type="checkbox"/> Fanconi anemia  | <input type="checkbox"/> Retinoblastoma, hereditary               |
| <input type="checkbox"/> Gastric cancer, hereditary diffuse                      | <input type="checkbox"/> Rhabdoid predisposition syndrome         |
| <input type="checkbox"/> Gastrointestinal stromal tumor, hereditary              | <input type="checkbox"/> Rothmund-Thomson syndrome                |
| <input type="checkbox"/> Hyperparathyroidism, familial                           | <input type="checkbox"/> Simpson-Golabi-Behmel syndrome           |
| <input type="checkbox"/> Leukemia, acute myeloid, familial                       | <input type="checkbox"/> Testicular germ cell tumor, familial     |
| <input type="checkbox"/> Leukemia, chronic lymphocytic, familial                 | <input type="checkbox"/> Thyroid cancer, familial medullary       |
| <input type="checkbox"/> Li-Fraumeni syndrome                                    | <input type="checkbox"/> Thyroid cancer, familial non-medullary   |
| <input type="checkbox"/> Lymphoma, Hodgkin, familial                             | <input type="checkbox"/> Tuberosus sclerosis complex              |
| <input type="checkbox"/> Lymphoma, non-Hodgkin, familial                         | <input type="checkbox"/> von Hippel-Lindau syndrome               |
| <input type="checkbox"/> Melanoma, hereditary, multiple                          | <input type="checkbox"/> Waardenstrom macroglobulinemia, familial |
| <input type="checkbox"/> Mosaic variegated aneuploidy                            | <input type="checkbox"/> Werner syndrome                          |
|  | <input type="checkbox"/> Wilms tumor, familial                    |
|  | <input type="checkbox"/> Xeroderma pigmentosum                    |
|  | <input type="checkbox"/> Select All                               |

**11. \* Please note your membership in any of the following national societies or special interest groups. (Choose one or more)**

- American College of Medical Genetics (ACMG)
- American Psychological Association (APA)
- American Society of Clinical Oncology (ASCO)
- American Society of Human Genetics (ASHG)
- Collaborative Group of the Americas on Inherited Colorectal Cancer (CGA-ICC)
- International Society for Gastrointestinal Hereditary Tumors (InSIGHT)
- International Society of Nurses in Genetics (ISONG)
- National Society of Genetic Counselors (NSGC)
- NSGC Special Interest Group in Cancer
- Oncology Nursing Society (ONS)
- ONS Cancer Genetics Special Interest Group

**12. Please click the Submit button to complete your application.**

We will process your application within a week and send you an email with a link to your listing on the Cancer.gov Web site. Please review the listing and let us know if any changes need to be made. You will also receive a verification request by email once a year.

For more information about the directory or help with the application, please contact the Directory Coordinator at [GeneticsDirectory@cancer.gov](mailto:GeneticsDirectory@cancer.gov).

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