



**ACGME International**

**Advanced Specialty Program Requirements for  
Graduate Medical Education in  
Medical Oncology  
(Internal Medicine)**

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# ACGME International Specialty Program Requirements for Graduate Medical Education in Medical Oncology (Internal Medicine)

## Int. Introduction

*Background and Intent: Programs must achieve and maintain Foundational Accreditation according to the ACGME-I Foundational Requirements prior to receiving Advanced Specialty Accreditation. The Advanced Specialty Requirements noted below complement the ACGME-I Foundational Requirements. For each section, the Advanced Specialty Requirements should be considered together with the Foundational Requirements.*

### Int. I. Definition and Scope of the Specialty

The specialty of medical oncology focuses on the etiology, diagnosis, prevention, and treatment of tumors (cancer)-benign and malignant neoplasms.

### Int. II. Duration of Education

Int. II.A. The educational program in medical oncology must be 24 or 36 months in length.

## I. Institution

### I.A. Sponsoring Institution

I.A.1. A fellowship in medical oncology must function as an integral part of an ACGME-I-accredited residency in internal medicine.

### I.B. Participating Sites

See International Foundational Requirements, Section I.B.

## II. Program Personnel and Resources

### II.A. Program Director

See International Foundational Requirements, Section II.A.

### II.B. Faculty

II.B.1. Qualified faculty members in the following subspecialties should be available for the education of the fellows:

II.B.1.a) cardiovascular disease;

II.B.1.b) endocrinology;

II.B.1.c) gastroenterology;

II.B.1.d) hospice and palliative medicine;

45	II.B.1.e)	<u>infectious diseases; and,</u>
46	II.B.1.f)	<u>pulmonary disease.</u>
47		
48	<b>II.C.</b>	<b>Other Program Personnel</b>
49		
50	II.C.1.	<u>The fellowship must have access to Clinical specialists,</u>
51		<u>including dermatologists, neurological surgeons, neurologists, obstetrician-</u>
52		<u>gynecologists, orthopaedic surgeons, otolaryngologists, radiation</u>
53		<u>oncologists, and urologists must participate in the education of fellows.</u>
54		
55	II.C.2.	<u>The fellowship must have access to surgeons in general surgery and</u>
56		<u>other surgical specialties, including those with a special interest in</u>
57		<u>oncology must participate in the education of fellows.</u>
58		
59	II.C.3.	Expertise in the following disciplines should be available to the program to
60		provide multidisciplinary patient care and fellow education:
61		
62	II.C.3.a)	genetic counseling;
63		
64	<del>II.C.3.b)</del>	<del>hospice and palliative care;</del>
65		
66	II.C.3.b)	oncologic nursing;
67		
68	II.C.3.c)	pain management;
69		
70	II.C.3.d)	psychiatry; and,
71		
72	II.C.3.e)	rehabilitation medicine.
73		
74	<b>II.D.</b>	<b>Resources</b>
75		
76	II.D.1.	Laboratory and imaging services must be available, including:
77		
78	II.D.1.a)	a hematology laboratory located at the primary clinical site; and,
79		
80	II.D.1.b)	a specialized coagulation laboratory.
81		
82	II.D.2.	Imaging services must be available, including:
83	II.D.2.a)	cross-sectional imaging, including computed tomography (CT) and
84		magnetic resonance imaging (MRI);
85	II.D.2.b)	nuclear medicine imaging; and,
86	II.D.2.c)	positron emission tomography (PET) scan imaging.
87		
88	II.D.3.	There must be advanced pathology services, including:
89		
90	II.D.3.a)	blood banking;
91		
92	II.D.3.b)	immunopathology; and,

93		
94	II.D.3.c)	transfusion and apheresis.
95		
96	II.D.4.	There must be a hematology clinical program with which fellows may
97		interact.
98	II.D.5.	Radiation oncology facilities must be available.
99		
100	<b>III.A.</b>	<b>Eligibility Criteria</b>
101		
102	III.A.1.	Prior to appointment in the program, fellows should have completed an
103		ACGME-I-accredited residency program in internal medicine, or an
104		internal medicine residency program acceptable to the Sponsoring
105		Institution's Graduate Medical Education Committee.
106		
107	<b>III.B.</b>	<b>Number of Fellows</b>
108		
109		See International Foundational Requirements, Section III.B.
110		
111	<b>IV.</b>	<b>Specialty-Specific Educational Program</b>
112		
113	<b>IV.A.</b>	<b>ACGME-I Competencies</b>
114	IV.A.1.	The program must integrate the following ACGME-I Competencies into the
115		curriculum.
116	IV.A.1.a)	Professionalism
117		
118	IV.A.1.a).(1)	Fellows must demonstrate a commitment to
119		professionalism and an adherence to ethical principles.
120		Fellows must demonstrate:
121		
122	IV.A.1.a).(1).(a)	personal development, attitudes, and coping skills of
123		physicians who care for critically ill patients.
124		
125	IV.A.1.b)	Patient Care and Procedural Skills
126		
127	IV.A.1.b).(1)	Fellows must provide patient care that is compassionate,
128		appropriate, and effective for the treatment of health
129		problems and the promotion of health. Fellows must
130		demonstrate competence in managing the care of
131		patients:
132		
133	IV.A.1.b).(1).(a)	<u>in a variety of health care settings, including</u>
134		<u>inpatient and ambulatory settings;the the</u>
135		<u>practice of health promotion, disease</u>
136		<u>prevention, diagnosis, care, and treatment of</u>
137		<u>patients of each gender, from adolescence to old</u>
138		<u>age, during health and all stages of illness;</u>
139	IV.A.1.b).(1).(b)	<u>using critical thinking and evidence-based tools;</u>

140	IV.A.1.b).(1).(c)	<u>using population-based data; and,</u>
141	IV.A.1.b).(1).(d)	<u>with whom they have limited or no physical contact,</u>
142		<u>through the use of telemedicine.</u>
143		
144	IV.A.1.b).(2).	<u>Fellows must demonstrate competence in</u> assuming
145		continuing responsibility for acutely and chronically ill patients
146		with medical oncology disorders in both inpatient and
147		outpatient settings, as well as the natural history of their
148		cancers, and the benefits and adverse effects of their
149		therapies.
150		
151	IV.A.1.b).(3).	<u>Fellows must demonstrate competence in</u> prevention,
152		evaluation, diagnosis, cancer staging, and management of
153		patients with <del>neoplastic</del> <u>malignant</u> disorders of the:
154	IV.A.1.b).(3).(a)	breast;
155		
156	IV.A.1.b).(3).(b)	cancer family syndromes;
157		
158	IV.A.1.b).(3).(c)	central nervous system;
159		
160	IV.A.1.b).(3).(d)	gastrointestinal tract (esophagus, stomach, colon,
161		rectum, anus);
162		
163	IV.A.1.b).(3).(e)	genitourinary tract;
164		
165	IV.A.1.b).(3).(f)	gynecologic malignancies;
166		
167	IV.A.1.b).(3).(g)	head and neck;
168		
169	IV.A.1.b).(3).(h)	hematopoietic system, <u>including myeloproliferative</u>
170		<u>neoplasms, myelodysplasias, acute and chronic</u>
171		<u>leukemias, Castleman disease, and dendritic cell</u>
172		<u>disorders;</u>
173		
174	IV.A.1.b).(3).(i)	liver;
175		
176	IV.A.1.b).(3).(j)	lung;
177		
178	IV.A.1.b).(3).(k)	lymphoid organs, <u>including lymphomas, myeloma, and</u>
179		<u>plasma cell dyscrasias;</u>
180		
181	IV.A.1.b).(3).(l)	pancreas;
182		
183	IV.A.1.b).(3).(m)	skin, including melanoma;
184		
185	IV.A.1.b).(3).(n)	testes; and,
186		
187	IV.A.1.b).(3).(o)	thyroid and other endocrine organs, including multiple
188		endocrine neoplasia <del>(MEN)</del> syndromes.
189		

190	IV.A.1.b).(4)	<u>Fellows must demonstrate competence in pathogenesis,</u>
191		<u>diagnosis, prevention, evaluation, and management of</u>
192		<u>patients with disorders whose characteristics overlap with</u>
193		<u>the areas of classical and malignant hematology, including:</u>
194	IV.A.1.b).(4).(a)	<u>bone marrow failure syndromes;</u>
195	IV.A.1.b).(4).(b)	<u>histiocytic disorders;</u>
196		
197	IV.A.1.b).(4).(c)	<u>myelodysplastic syndromes; and,</u>
198	IV.A.1.b).(4).(d)	<u>myeloproliferative neoplasms.</u>
199	IV.A.1.b).(5)	<u>Fellows must demonstrate competence in the diagnosis and</u>
200		<u>management of classical hematologic complications of</u>
201		<u>malignant disorders, including:</u>
202	IV.A.1.b).(5).(a)	<u>autoimmune disorders, to include hemolytic anemia</u>
203		<u>and other hematologic manifestations of autoimmune</u>
204		<u>disorders;</u>
205	IV.A.1.b).(5).(b)	<u>congenital and acquired thrombotic disorders;</u>
206	IV.A.1.b).(5).(c)	<u>hemoglobin disorders, to include sickle cell disease</u>
207		<u>and thalassemia syndromes;</u>
208	IV.A.1.b).(5).(d)	<u>hemophilias, von Willebrand disease, and other</u>
209		<u>inherited and acquired hemorrhagic disorders, to</u>
210		<u>include platelet function defects;</u>
211		
212	IV.A.1.b).(5).(e)	<u>inherited and acquired disorders of the red blood cell</u>
213		<u>membrane and of red blood cell metabolism;</u>
214	IV.A.1.b).(5).(f)	<u>inherited and acquired disorders of white blood cells;</u>
215	IV.A.1.b).(5).(g)	<u>nutritional anemias;</u>
216	IV.A.1.b).(5).(h)	<u>platelet disorders, including idiopathic</u>
217		<u>thrombocytopenic purpura(ITP) and congenital</u>
218		<u>thrombocytopenias;</u>
219	IV.A.1.b).(5).(i)	<u>the porphyrias; and,</u>
220	IV.A.1.b).(5).(j)	<u>thrombotic microangiopathies.</u>
221	IV.A.1.b).(6)	<u>Fellows must be able to perform all medical, diagnostic, and</u>
222		<u>surgical procedures considered essential to the subspecialty,</u>
223		<u>including:</u>
224	IV.A.1.b).(6).(a)	<u>performing diagnostic and therapeutic procedures</u>
225		<u>relevant to their specific career path, to include care</u>
226		<u>and management of venous access devices.</u>

227	IV.A.1.b).(6).(b)	<u>treating their patients' conditions with practices that are</u>
228		<u>patient-centered, safe, scientifically based, effective,</u>
229		<u>timely, and cost-effective, including:</u>
230	IV.A.1.b).(6).(b).(i)	care and management of the geriatric patient
231		with malignancy and hematologic disorders,
232		to <u>include Castleman disease;</u>
233		
234	IV.A.1.b).(6).(b).(ii)	care of patients with human immunodeficiency
235		virus(HIV)-related malignancies;
236	IV.A.1.b).(6).(b).(ii)	<u>hematologic care of pregnant patients and</u>
237		<u>women of reproductive age;</u>
238	IV.A.1.b).(6).(b).(iii)	<u>hematologic care of transgender patients;</u>
239	IV.A.1.b).(6).(b).(iv)	<u>hematologic complications of infectious</u>
240		<u>diseases;</u>
241		
242	IV.A.1.b).(6).(b).(v)	management of pain, anxiety, and depression
243		in patients with cancer;
244		
245	IV.A.1.b).(6).(b).(vi)	management of the neutropenic and the
246		immunocompromised patient;
247		
248	IV.A.1.b).(6).(b).(vii)	palliative care, to include hospice and home
249		care;
250		
251	IV.A.1.b).(6).(b).(viii)	rehabilitation and psychosocial care of patients
252		with cancer;
253		
254	IV.A.1.b).(6).(b).(ix)	specific cancer prevention and screening for
255		high-risk individuals, to include genetic
256		testing;
257		
258	IV.A.1.b).(6).(b).(x)	treatment and diagnosis of recognition and
259		management of paraneoplastic disorders;
260		
261	IV.A.1.b).(6).(b).(xi)	use of chemotherapeutic agents and biological
262		products through all therapeutic routes;
263		
264	IV.A.1.b).(6).(b).(xii)	use of chemotherapeutic drugs, biologic
265		products, and growth factors; their
266		mechanisms of action, pharmacokinetics,
267		clinical indications, and limitations; to include
268		their effects, toxicity, and interactions;
269		
270	IV.A.1.b).(6).(b).(xiii)	use of hematologic, infectious disease, and
271		nutrition support;
272	IV.A.1.b).(6).(b).(xiv)	<u>use of immunotherapeutic drugs; their</u>
273		<u>mechanisms of action, pharmacokinetics,</u>

274		<u>clinical indications, and limitations; and their</u>
275		<u>effects, toxicity, and interactions; to include</u>
276		<u>the use of cellular immunotherapies (such as,</u>
277		<u>CAR-T therapies);</u>
278		
279	IV.A.1.b).(6).(b).(xv)	use of multi-agent chemotherapeutic protocols
280		and combined modality therapy of neoplastic
281		disorders; and,
282	IV.A.1.b).(6).(b).(xvi)	<u>use of systemic therapies through all</u>
283		<u>therapeutic routes.</u>
284	IV.A.1. b).(6).(c)	<u>using diagnostic and/or imaging studies relevant to the</u>
285		<u>care of the patient, including:</u>
286		
287	IV.A.1.b).(6).(c).(i)	assessment of tumor burden (and response
288		as measured by physical and radiologic exam)
289		and tumor markers;
290		
291	IV.A.1.b).(6).(c).(ii)	assessment of tumor imaging by CT, MRI, PET
292		scanning, and nuclear imaging techniques;
293	IV.A.1.b).(6).(c).(iii)	correlation of clinical information with cytology,
294		histology, and immunodiagnostic imaging
295		techniques; and,
296	IV.A.1.b).(6).(c).(iv)	<u>indications and application of imaging</u>
297		<u>techniques in patients with neoplastic</u>
298		<u>disorders.</u>
299		
300	IV.A.1.c)	Medical Knowledge
301	IV.A.1.c).(1)	Fellows must demonstrate knowledge of established and
302		evolving biomedical clinical, epidemiological, and social-
303		behavioral sciences, as well as the application of this
304		knowledge to patient care. Fellows must demonstrate
305		knowledge of:
306		
307	IV.A.1.c).(1).(a)	the scientific method of problem solving and
308		evidence-based decision-making;
309		
310	IV.A.1.c).(1).(b)	indications, contraindications, and techniques for,
311		and limitations, complications, and interpretation of
312		results of those diagnostic and therapeutic
313		procedures integral to the discipline, including the
314		appropriate indications for and use of screening
315		tests and procedures;
316		
317	IV.A.1.c).(1).(c)	basic molecular and pathophysiologic
318		mechanisms, diagnosis, and therapy of diseases of
319		the blood, to include anemias, diseases of white
320		blood cells and stem cells, and disorders of



321		hemostasis and thrombosis;
322	IV.A.1.c).(1).(d)	clinical epidemiology and biostatistics, including
323		clinical study and experimental protocol design, data
324		collection, and analysis;
325	IV.A.1.c).(1).(e)	<u>functional characteristics, indications, risks, and</u>
326		<u>process of using indwelling venous access devices.</u>
327	IV.A.1.c).(1).(f)	genetics and developmental biology, including;
328	IV.A.1.c).(1).(f).(i)	cytogenetics;
329	IV.A.1.c).(1).(f).(ii)	molecular genetics; and,
330	IV.A.1.c).(1).(f).(iii)	the nature of oncogenes and their products.
331	IV.A.1.c).(1).(g)	gene therapy;
332	IV.A.1.c).(1).(h)	immune markers, immunophenotyping, flow
333		cytometry, cytochemical studies, and cytogenetic
334		and DNA analysis of neoplastic disorders;
335		
336	IV.A.1.c).(1).(i)	indications for, complications of, and risks and
337		limitations associated with:
338		
339	IV.A.1.c).(1).(i).(i)	lesion biopsy <u>detection of circulating DNA for</u>
340		<u>disease-specific markers;</u>
341		
342	IV.A.1.c).(1).(m).(ii)	lumbar puncture;
343		
344	IV.A.1.c).(1).(i).(ii)	paracentesis;
345		
346	IV.A.1.c).(1).(i).(iii)	skin biopsies; and,
347	IV.A.1.c).(1).(i).(iv)	thoracentesis.
348		
349	IV.A.1.c).(1).(j)	malignant and hematologic complications of
350		organ transplantation;
351		
352	IV.A.1.c).(1).(k)	management of post-transplant complications;
353		
354	IV.A.1.c).(1).(l)	mechanisms of action, pharmacokinetics, clinical
355		indications for, and limitations of chemotherapeutic
356		drugs, and biologic products <u>including cellular</u>
357		<u>immunotherapies (such as CAR-T therapies); and</u>
358		<u>growth factors</u> , including their effects, toxicity, and
359		interactions,
360	IV.A.1.c).(1).(m)	pathogenesis, diagnosis, and treatment of disease,
361		including etiology, epidemiology, natural history,
362		diagnosis, pathology, staging, and management

363		of neoplastic diseases of the blood, blood-forming
364		organs, and lymphatic tissues.
365	IV.A.1.c).(1).(n)	physiology and pathophysiology, including:
366	IV.A.1.c).(1).(n).(i)	basic and clinical pharmacology,
367		pharmacokinetics, and toxicity;
368		
369	IV.A.1.c).(1).(n).(ii)	cell and molecular biology;
370		
371	IV.A.1.c).(1).(n).(iii)	hematopoiesis;
372		
373	IV.A.1.c).(1).(n).(iv)	molecular mechanisms of hematopoietic and
374		lymphopoietic malignancies;
375	IV.A.1.c).(1).(n).(v)	pathophysiology and patterns of tumor
376		metastases;
377		
378	IV.A.1.c).(1).(n).(vi)	principles of oncogenesis; and,
379		
380	IV.A.1.c).(1).(n).(vii)	tumor immunology.
381		
382	IV.A.1.c).(1).(o)	principles of, indications for, and complications of
383		<del>autologous and allogeneic bone marrow or</del>
384		peripheral blood stem cell transplantation;
385		
386	IV.A.1.c).(1).(p)	principles of, indications for, and complications of
387		peripheral stem cell harvests;
388		
389	IV.A.1.c).(1).(q)	principles of, indications for, and limitations of:
390	IV.A.1.c).(1).(q).(i)	surgery in the treatment of cancer; and,
391	IV.A.1.c).(1).(q).(ii)	radiation therapy in the treatment of cancer.
392		
393	IV.A.1.c).(1).(r)	the basic principles of laboratory and clinical testing,
394		quality control, quality assurance, and proficiency
395		standards.
396	IV.A.1.c).(2)	<u>Fellows must demonstrate sufficient knowledge specific</u>
397		<u>to the subspecialty of medical oncology, including</u>
398		<u>application of technology appropriate for the clinical</u>
399		<u>context, to include evolving technologies.</u>
400		
401	IV.A.1.d)	Practice-Based Learning and Improvement
402		
403	IV.A.1.d).(1)	Fellows must demonstrate the ability to investigate and
404		evaluate their care of patients, to appraise and assimilate
405		scientific evidence, and to continuously improve patient
406		care based on constant self-evaluation and lifelong
407		learning.

408	IV.A.1.e)	Interpersonal and Communication Skills
409	IV.A.1.e).(1)	Fellows must demonstrate interpersonal and communication
410		skills that result in the effective exchange of information and
411		collaboration with patients, patients' families, and health
412		professionals. Fellows must demonstrate:
413		
414	IV.A.1.e).(1).(a)	team leadership skills and the ability to work with an
415		interdisciplinary team by:
416		
417	IV.A.1.e).(1).(a).(i)	identifying essential team members;
418		
419	IV.A.1.e).(1).(a).(ii)	defining the roles of team members; and,
420		
421	IV.A.1.e).(1).(a).(iii)	evaluating the role of the interdisciplinary
422		team.
423		
424	IV.A.1.f)	Systems-Based Practice
425		
426	IV.A.1.f).(1)	Fellows must demonstrate an awareness of and
427		responsiveness to the larger context and system of health
428		care, including the social determinates of health, as well as
429		the ability to call effectively on other resources in the
430		system to produce optimal care.
431		
432	<b>IV.B.</b>	<b>Regularly Scheduled Educational Activities</b>
433	IV.B.1.	<u>The educational program must include didactic instruction based on the</u>
434		<u>core knowledge content in medical oncology.</u>
435	IV.B.1.b)	<u>Fellows must have a sufficient number of didactic sessions to</u>
436		<u>ensure fellow-fellow and fellow-and-faculty member interaction.</u>
437	IV.B.2.	<u>The program must ensure that fellows have an opportunity to review all</u>
438		<u>knowledge content from conferences that they could not attend.</u>
439	IV.B.3.	Fellows must participate in multidisciplinary case management or tumor
440		board conferences and in protocol studies.
441	IV.B.4.	<u>Fellows must receive instruction in practice management relevant to</u>
442		<u>medical oncology.</u>
443	<b>IV.C.</b>	<b>Clinical Experiences</b>
444	IV.C.1.	<u>Assignment of rotations must be structured to minimize the frequency of</u>
445		<u>rotational transitions, and rotations must be of sufficient length to provide a</u>
446		<u>quality educational experience, defined by continuity of patient care, ongoing</u>
447		<u>supervision, longitudinal relationships with faculty members, and meaningful</u>
448		<u>assessment and feedback.</u>
449	IV.C.2.	<u>Rotations must be structured to allow fellows to function as a part of an</u>

450		<u>effective interprofessional team that works together toward the shared goals</u>
451		<u>of patient safety and quality improvement.</u>
452	IV.C.3.	<u>Rotations must be structured to minimize conflicting inpatient and outpatient</u>
453		<u>responsibilities.</u>
454		
455	IV.C.4.	At least 12 months of education must be devoted to clinical experience.
456	IV.C.4.a)	At least 50 percent of the clinical experience must occur in the
457		outpatient setting.
458		
459	IV.C.4.b)	<del>The program must provide at least one month of clinical</del>
460		<del>experience in autologous bone marrow transplantation.</del>
461		
462	IV.C.5.	Inpatient assignments should be of sufficient duration to permit continuing
463		care of a majority of a fellow's patients throughout their hospitalization.
464	IV.C.6.	<u>The program must provide educational experiences in team-based care</u>
465		<u>that allow fellows to interact with and learn from other health care</u>
466		<u>professionals.</u>
467	IV.C.7.	<u>The educational program must provide fellows with elective experiences</u>
468		<u>relevant to their future practice or to further skill/competence development.</u>
469	IV.C.7.a)	<u>Fellows should have the opportunity to develop competence in</u>
470		<u>performing thoracentesis, paracentesis, and skin and lesion</u>
471		<u>biopsies.</u>
472		
473	IV.C.7.b)	<u>Additional training and experiences should be made available for</u>
474		<u>those fellows who request the need to perform specified</u>
475		<u>procedures in their post-fellowship careers (such as bone</u>
476		<u>marrow aspirates, lumbar punctures for diagnosis and/or</u>
477		<u>administration of intrathecal chemotherapy, administering</u>
478		<u>therapeutics through Ommaya reservoirs).</u>
479	IV.C.8.	Fellows must participate in training using simulation.
480		
481	IV.C.9.	Fellows should have a structured continuity ambulatory clinic experience
482		<u>for the duration of the program</u> that exposes them to the breadth and
483		depth of medical oncology.
484	IV.C.9.a)	<del>This should include an appropriate distribution of patients of each</del>
485		<del>gender and a diversity of ages.</del>
486		
487	IV.C.9.a)	The experience should average one half-day each week
488		throughout the education program.
489		
490	IV.C.9.a).(1)	<del>Each fellow should, on average, be responsible for four</del>
491		<del>to eight patients during each half-day session.</del>
492		
493	IV.C.9.a).(1).(a)	<del>Each fellow should, on average, be responsible</del>
494		<del>for no more than eight to 12 patients during</del>

each half day ambulatory session.

- IV.C.9.b) The continuing patient care experience should not be interrupted by more than one month, excluding a fellow's vacation.

#### **IV.D. Scholarly Activity**

##### **IV.D.1. Fellows' Scholarly Activity**

- IV.D.1.a) While in the program, each fellow must complete at least one of the following scholarly activities: participation in grand rounds; poster presentations; workshops; quality improvement presentations; podium presentations; grant leadership; non-peer-reviewed print/electronic resources; articles or publications; book chapters; textbooks; webinars; service on professional committees; or service as a journal reviewer, journal editorial board member, or editor.

##### **IV.D.2. Faculty Scholarly Activity**

See International Foundational Requirements, Section IV.D.2.

#### **V. Evaluation**

See International Foundational Requirements, Section V.

#### **VI. The Learning and Working Environment**

##### **VI.A. Principles**

See International Foundational Requirements, Section VI.A.

##### **VI.B. Patient Safety**

See International Foundational Requirements, Section VI.B.

##### **VI.C. Quality Improvement**

See International Foundational Requirements, Section VI.C.

##### **VI.D. Supervision and Accountability**

- VI.D.1. Direct supervision of procedures performed by each fellow must occur until competence has been acquired and documented by the program director.

##### **VI.E. Professionalism**

See International Foundational Requirements, Section VI.E.

##### **VI.F. Well-being**

542		See International Foundational Requirements, Section VI.F.
543		
544	<b>VI.G.</b>	<b>Fatigue</b>
545		
546		See International Foundational Requirements, Section VI.G.
547		
548	<b>VI.H.</b>	<b>Transitions of Care</b>
549		
550		See International Foundational Requirements, Section VI.H.
551		
552	<b>VI.I.</b>	<b>Clinical Experience and Education</b>
553		
554		See International Foundational Requirements, Section VI.I.
555		
556	<b>VI.J.</b>	<b>On-Call Activities</b>
557		
558		See International Foundational Requirements, Section VI.J.