Department of Veterans Affairs	HEMATOLOGIC AND LYMPHATIC CONDITIONS, INCLUDING LEUKEMIA DISABILITY BENEFITS QUESTIONNAIRE		
NAME OF PATIENT/VETERAN		PATIENT/VETERAN'S SOCIAL SECU	IRITY NUMBER
IMPORTANT - THE DEPARTMENT OF VETERANS AFFA COMPLETING AND/OR SUBMITTING THIS FORM.	IRS (VA) WILL NOT PAY OR REIMBURSE A	NY EXPENSES OR COST INCURRED IN THE PRO	OCESS OF
Note - The Veteran is applying to the U.S. Department of V of their evaluation in processing the Veteran's claim. VA m veteran's application. VA reserves the right to confirm the a by the Veteran's provider .	ay obtain additional medical information, inclu	ding an examination, if necessary, to complete VA's	review of the
Are you completing this Disability Benefits Questionnaire	at the request of:		
Veteran/Claimant			
Other: please describe			
Are you a VA Healthcare provider? O Yes O No)		
Is the Veteran regularly seen as a patient in your clinic?	Yes No		
Was the Veteran examined in person? O Yes	No		
If no, how was the examination conducted?			
	EVIDENCE REVIEW		
Evidence reviewed:			
○ No records were reviewed			
C Records reviewed			
Please identify the evidence reviewed (e.g. service treatm	nent records, VA treatment records, private tre	atment records) and the date range.	

SE	SECTION I - DIAGNOSIS				
1A. CHECK THE CLAIMED HEMATOLOGICAL AND/OR LYMPHATIC CON	DITION(S) THAT PER	TAIN TO THIS DBQ:			
NOTE: These are the diagnoses determined during this current evaluation of from a previous diagnosis for this condition, or if there is a diagnosis of a con section. Date of diagnosis can be the date of evaluation if the clinician is mak history.	nplication due to the cla	aimed condition, explain your findings and reasons in the comments			
Agranulocytosis, acquired		DATE OF DIAGNOSIS:			
Chronic myelogenous leukemia (CML) (chronic myeloid leukemia chronic granulocytic leukemia)	^{I Or} ICD CODE:	DATE OF DIAGNOSIS:			
Chronic lymphocytic leukemia (CLL)	ICD CODE:	DATE OF DIAGNOSIS:			
Hairy cell or other B-cell leukemia	ICD CODE:	DATE OF DIAGNOSIS:			
Other	ICD CODE:	DATE OF DIAGNOSIS:			
Hodgkin's lymphoma Active disease Treatment phase	ICD CODE:	DATE OF DIAGNOSIS:			
Non-Hodgkin's lymphoma		DATE OF DIAGNOSIS:			
Active disease Treatment phase Indolent and	non-contiguous phase	e of low grade NHL			
Multiple myeloma	ICD CODE:	DATE OF DIAGNOSIS:			
Myelodysplastic syndrome	ICD CODE:	DATE OF DIAGNOSIS:			
Solitary plasmacytoma	ICD CODE:	DATE OF DIAGNOSIS:			
Anemia					
Aplastic anemia	ICD CODE:	DATE OF DIAGNOSIS:			
Iron deficiency anemia		DATE OF DIAGNOSIS:			
Folic acid deficiency	ICD CODE:	DATE OF DIAGNOSIS:			
Pernicious anemia or other Vitamin B12 deficiency anemia		DATE OF DIAGNOSIS:			
Acquired hemolytic anemia		DATE OF DIAGNOSIS:			
AL amyloidosis (primary amyloidosis)					
Immune thrombocytopenia					
Polycythemia vera	ICD CODE:				
Sickle cell anemia	ICD CODE:	DATE OF DIAGNOSIS:			
Splenectomy	ICD CODE:	DATE OF DIAGNOSIS:			
Are there complications such as systemic infections with encapsulated	bacteria?	ES 🗌 NO			
If Yes, complete SECTION VIII - OTHER PERTINENT PHYSICAL FIN					
linium to Splace					
Injury to Spleen If checked, complete SECTION VIII - OTHER PERTINENT PHYSICAL					
Adenitis, tuberculous (Also complete the Infectious Diseases (Other Than HIV-Related Illness, Chronic Fatigue Syndrome, or Tuberculosis) Disability Benefits Questionnaire).		DATE OF DIAGNOSIS:			
Active Inactive					
Essential thrombocythemia or primary myelofibrosis	ICD CODE:	DATE OF DIAGNOSIS:			
Other, specify					
Other diagnosis #1:	ICD CODE:	DATE OF DIAGNOSIS:			
Other diagnosis #2:	ICD CODE:	DATE OF DIAGNOSIS:			
Other diagnosis #3:	ICD CODE:	DATE OF DIAGNOSIS:			
1B. IF THERE ARE ADDITIONAL OR PRIOR DIAGNOSES THAT PERTAIN	TO HEMATOLOGIC	OR LYMPHATIC CONDITIONS, LIST USING ABOVE FORMAT:			
Llemetalegie and Lymphotic Conditions Dischility Deposite Ouestiann		Undated on March 31, 2020 -v/20			

SECTION II - MEDICAL HISTORY		
2A. DESCRIBE THE HISTORY (including cause (if known), onset and course) OF THE VETERAN'S CURRENT HEMATOLOGIC OR LYMPHATIC CONDITION(S) (brief summary):		
2B. IS CONTINUOUS MEDICATION REQUIRED FOR CONTROL OF A HEMATOLOGIC OR LYMPHATIC CONDITION, INCLUDING ANEMIA OR THROMBOCYTOPENIA CAUSED BY TREATMENT FOR A HEMATOLOGIC OR LYMPHATIC CONDITION?		
YES NO		
IF YES, LIST ONLY THOSE MEDICATIONS REQUIRED FOR CONTROL OF THE VETERAN'S HEMATOLOGIC OR LYMPHATIC CONDITION, INCLUDING ANEMIA OR THROMBOCYTOPENIA CAUSED BY TREATMENT FOR A HEMATOLOGIC OR LYMPHATIC CONDITION. PROVIDE THE NAME OF THE MEDICATION AND THE CONDITION THE MEDICATION IS USED TO TREAT:		
2C. INDICATE THE STATUS OF THE PRIMARY HEMATOLOGIC OR LYMPHATIC CONDITION:		
SECTION III - TREATMENT		
3A. HAS THE VETERAN COMPLETED ANY TREATMENT OR IS THE VETERAN CURRENTLY UNDERGOING ANY TREATMENT FOR ANY HEMATOLOGIC OR LYMPHATIC CONDITION, INCLUDING LEUKEMIA?		
YES NO; WATCHFUL WAITING		
IF YES, INDICATE TYPE OF TREATMENT THE VETERAN IS CURRENTLY UNDERGOING OR HAS COMPLETED (Check all that apply):		
 Treatment completed; currently in watchful waiting status Transplant (specify type) 		
Peripheral blood stem cell transplant Bone marrow stem cell transplant Other (specify)		
If checked, provide:		
Date of hospital admission and location: Date of hospital discharge after transplant:		
Surgery, if checked describe:		
Date(s) of surgery:		
Radiation therapy		
Date of most recent treatment:		
Date of completion of treatment or anticipated date of completion:		
Antineoplastic chemotherapy		
Date of most recent treatment:		
Date of completion of treatment or anticipated date of completion:		
Other therapeutic procedure		
If checked, describe procedure:		
Other therapeutic treatment If checked, describe treatment:		
Date of completion of treatment or anticipated date of completion:		
SECTION IV - ANEMIA AND THROMBOCYTOPENIA		
4A. DOES THE VETERAN HAVE ANEMIA OR THROMBOCYTOPENIA, INCLUDING THAT CAUSED BY TREATMENT FOR A HEMATOLOGIC OR		
YES NO		
IF YES, COMPLETE THE FOLLOWING:		

SECTION IV - ANEMIA AND THROMBOCYTOPENIA (Continued)
4B. DOES THE VETERAN HAVE ANEMIA (other than Sickle Cell Anemia) OR THROMBOCYTOPENIA?
YES NO
IF YES, PLEASE CHECK TYPE:
 Aplastic anemia (complete 4C) Iron deficiency anemia (complete 4D) Folic acid deficiency (complete 4E) Pernicious anemia or other Vitamin B12 deficiency anemia (complete 4F) Acquired hemolytic anemia (complete 4G) Immune thrombocytopenia (complete 4H) Other, specify
IS THE ANEMIA CAUSED BY TREATMENT FOR ANOTHER HEMATOLOGIC OR LYMPHATIC CONDITION?
YES NO
IF YES, PROVIDE THE NAME OF THE OTHER HEMATOLOGIC OR LYMPHATIC CONDITION CAUSING THE SECONDARY ANEMIA:
4C. APLASTIC ANEMIA:
Requiring peripheral blood stem cell transplant
Requiring bone marrow stem cell transplant
Requiring transfusion of platelets, on average, at least: once every six weeks per 12-month period once every three months per 12-month period once per 12-month period
Requiring transfusion of red cells, on average, at least: once every six weeks per 12-month period once every three months per 12-month period once per 12-month period
Infections recurring, on average, at least: once every six weeks per 12-month period once every three months per 12-month period once per 12-month period
Using continuous therapy with immunosuppressive agent
Using continuous therapy with newer platelet stimulating factors
NOTE: The term "newer platelet stimulating factors" includes medication, factors, or other agents approved by the United States Food and Drug Administration.
4D. IRON DEFICIENCY ANEMIA
 Requiring intravenous iron infusions 4 or more times per 12-month period Requiring intravenous iron infusions at least 1 time but less than 4 times per 12-month period Requiring continuous treatment with oral supplementation Requiring treatment only by dietary modification Asymptomatic
4E. FOLIC ACID DEFICIENCY
Requiring continuous treatment with high-dose oral supplementation
Requiring treatment only by dietary modification Asymptomatic
Hematologic and Lymphatic Conditions Disability Benefits Questionnaire

SECTION IV - ANEMIA AND THROMBOCYTOPENIA (Continued)
4F. PERNICIOUS ANEMIA OR OTHER VITAMIN B12 DEFICIENCY ANEMIA
For initial diagnosis requiring transfusion due to severe anemia
If checked, provide the date of initial diagnosis requiring transfusion and
the date of hospital discharge or cessation of parenteral B12 therapy
Signs or symptoms related to central nervous system impairment, such as encephalopathy, myelopathy, or severe peripheral neuropathy, requiring
parenteral B12 therapy
Requiring continuous treatment with Vitamin B12 injections
Requiring continuous treatment with Vitamin B12 sublingual tablets
Requiring continuous treatment with high-dose oral tablets
Requiring continuous treatment with Vitamin B12 nasal spray or gel
NOTE: If there are any residual effects of pernicious anemia, such as neurologic involvement causing peripheral neuropathy, myelopathy, dementia, or related gastrointestinal residuals, ALSO complete appropriate Questionnaire for each condition.
4G. ACQUIRED HEMOLYTIC ANEMIA
Required a bone marrow transplant
Requiring continuous intravenous or immunosuppressive therapy (e.g., prednisone, Cytoxan, azathioprine, or rituximab)
Requiring immunosuppressive medication 4 or more times per 12-month period
Requiring 2-3 courses of immunosuppressive therapy per 12-month period
Requiring one course of immunosuppressive therapy per 12-month period
Asymptomatic
4H. IMMUNE THROMBOCYTOPENIA
Requiring chemotherapy for chronic refractory thrombocytopenia
Requiring immunosuppressive therapy
Platelet count 30,000 or below despite treatment
Platelet count higher than 30,000 but not higher than 50,000 with history of hospitalization because of severe bleeding requiring intravenous immune globulin, high dose parenteral corticosteroids, and platelet transfusions
Platelet count higher than 30,000 but not higher than 50,000 with mild mucous membrane bleeding which requires oral corticosteroid therapy or intravenous
Platelet count higher than 30,000 but not higher than 50,000 with immune thrombocytopenia which requires oral corticosteroid therapy or intravenous immune globulin
Platelet count higher than 30,000 but not higher than 50,000, not requiring treatment
Platelet count above 50,000 and asymptomatic
In remission
SECTION V - LEUKEMIA, MULTIPLE MYELOMA, AGRANULOCYTOSIS, ACQUIRED, ESSENTIAL THROMBOCYTHEMIA, PRIMARY MYELOFIBROSIS, AND MYELODYSPLASTIC SYNDROMES
5A. DOES THE VETERAN HAVE LEUKEMIA, MULTIPLE MYELOMA, AGRANULOCYTOSIS, ACQUIRED, ESSENTIAL THROMBOCYTHEMIA, PRIMARY MYELOFIBROSIS, OR MYELODYSPLASTIC SYNDROMES?
YES NO
IF YES, PLEASE CHECK TYPE:
Chronic lymphocytic leukemia (complete 5B)
Monoclonal B-cell lymphocytosis (MBL) (complete 5B)
Hairy cell or other B-cell leukemia (complete 5B)
Chronic myelogenous leukemia (complete 5B)
Chronic myeloid leukemia (complete 5B)
Chronic granulocytic leukemia (complete 5B)
Multiple myeloma (complete 5C)
Agranulocytosis, acquired (complete 5D)
Essential thrombocythemia or primary myelofibrosis (complete 5E)
Myelodysplastic syndromes (complete 5F)
Other, specify

SECTION V - LEUKEMIA, MULTIPLE MYELOMA, AGRANULOCYTOSIS, ACQUIRED, ESSENTIAL THROMBOCYTHEMIA, PRIMARY MYELOFIBROSIS, AND MYELODYSPLASTIC SYNDROMES (Continued)			
5B. WHAT IS THE STATUS OF LEUKEMIA?			
 Asymptomatic, Rai Stage 0 Requiring peripheral blood stem cell transplant Requiring bone marrow stem cell transplant Requiring continuous myelosuppressive therapy Requiring continuous immunosuppressive therapy treatment Requiring intermittent myelosuppressive therapy, or molecularly targeted therapy with tyrosine kinase inhibitors, or interferon treatment when not in apparent remission In apparent remission on continuous molecularly targeted therapy with tyrosine kinase inhibitors 			
5C. WHAT IS THE STATUS OF MULTIPLE MYELOMA?			
 Symptomatic (if checked, provide date of the diagnosis of symptomatic multiple myeloma) Asymptomatic Smoldering or monoclonal gammopathy of undetermined significance (MGUS) NOTE: Current validated biomarkers of symptomatic multiple myeloma, asymptomatic, smoldering or monoclonal gammopathy of undetermined significance (MGUS) are acceptable for the diagnosis of multiple myeloma as defined by the American Society of Hematology (ASH) and International Myeloma Working Group (IMWG). 			
5D. WHAT IS THE STATUS OF AGRANULOCYTOSIS, ACQUIRED?			
 Requiring bone marrow transplant Requiring intermittent myeloid growth factors (granulocyte colony-stimulating factor (G-CSF) or granulocyte-macrophage colony-stimulating factor (GM-CSF)) Requiring continuous immunosuppressive therapy such as cyclosporine to maintain absolute neutrophil count (ANC) greater than 500/microliter (µl) but less than 1000/µl Requiring intermittent myeloid growth factors to maintain ANC greater than 1000/µl Requiring intermittent use of a myeloid growth factor to maintain ANC greater than or equal to 1500/µl Infections recurring, on average, at least once every six weeks per 12-month period Infections recurring, on average, at least once per 12-month period but less than once every three months per 12-month period Requiring continuous medication (e.g., antibiotics) for control 			
5E. WHAT IS THE STATUS OF ESSENTIAL THROMBOCYTHEMIA AND PRIMARY MYELOFIBROSIS?			
Requiring continuous myelosuppressive therapy Requiring intermittent myelosuppressive therapy Requiring peripheral blood stem cell transplant Requiring bone marrow stem cell transplant Requiring chemotherapy Requiring interferon treatment Requiring interferon treatment to maintain platelet count < 500 x 10 ⁹ /L Requiring interferon treatment to maintain platelet count of 200,000-400,000 Requiring interferon treatment to maintain white blood cell (WBC) count of 4,000-10,000 Asymptomatic			
5F. WHAT IS THE STATUS OF MYELODYSPLASTIC SYNDROMES?			
 Requiring peripheral blood stem cell transplant Requiring bone marrow stem cell transplant Requiring chemotherapy Requiring 4 or more blood or platelet transfusions per 12-month period Requiring 1 to 3 blood or platelet transfusions per 12-month period Infections requiring hospitalization 3 or more times per 12-month period Infections requiring hospitalization 1 to 2 times per 12-month period Requiring biologic therapy on an ongoing basis Requiring erythropoiesis stimulating agent (ESA) for 12 weeks or less per 12-month period 			

6A. DOES THE VETERAN HAVE POLYCYTHEMIA VERA?
YES NO
IF YES, CHECK ALL THAT APPLY:
 Requiring peripheral blood or bone marrow stem-cell transplant for the purpose of ameliorating the symptom burden Requiring chemotherapy (including myelosuppressants) for the purpose of ameliorating the symptom burden Requiring phlebotomy 6 or more times per 12-month period or molecularly targeted therapy for the purpose of controlling RBC count Requiring phlebotomy 4-5 times per 12-month period to maintain platelets < 200,000 or white blood cells (WBC) < 12,000 Requiring phlebotomy 3 or fewer times per 12-month period to maintain all blood values at reference range levels Requiring continuous biologic therapy or myelosuppresive agents, to include interferon, to maintain platelets < 200,000 or white blood cells (WBC) < 12,000 Requiring biologic therapy or interferon on an intermittent basis as needed to maintain all blood values at reference range levels Other, describe:
NOTE: If there are complications due to polycythemia vera such as hypertension, gout, stroke or thrombotic disease, ALSO complete appropriate Questionnaire for each condition.
SECTION VII - SICKLE CELL ANEMIA
7A. DOES THE VETERAN HAVE SICKLE CELL ANEMIA?
IF YES, CHECK ALL THAT APPLY:
 Symptoms preclude even light manual labor Symptoms preclude other than light manual labor With anemia, thrombosis, and infarction With at least 4 or more painful episodes per 12-month period, occurring in skin, joints, bones, or any major organs caused by hemolysis and sickling of red blood cells With 3 painful episodes per 12-month period With 1 or 2 painful episodes per 12-month period With identifiable organ impairment In remission Asymptomatic Other, describe:
SECTION VIII - OTHER PERTINENT PHYSICAL FINDINGS, COMPLICATIONS, CONDITIONS, SIGNS AND/OR SYMPTOMS
BA. DOES THE VETERAN HAVE ANY OTHER PERTINENT PHYSICAL FINDINGS, COMPLICATIONS, CONDITIONS, SIGNS AND/OR SYMPTOMS RELATED TO THE CONDITIONS LISTED IN THE DIAGNOSIS SECTION ABOVE? YES NO IF YES, DESCRIBE (brief summary): Also if indicated, complete the appropriate questionnaire for each condition
8B. DOES THE VETERAN HAVE ANY SCARS OR OTHER DISFIGUREMENT (of the skin) RELATED TO ANY CONDITIONS OR TO THE TREATMENT OF ANY CONDITIONS LISTED IN THE DIAGNOSIS SECTION?
IF YES, ALSO COMPLETE APPROPRIATE DERMATOLOGICAL DBQ

SECTION IX - DIAGNOSTIC TESTING			
NOTE: If testing has been performed and reflects Veteran's current condition, no further testing is required. When appropriate, provide most recent complete blood count.			
9A. HAS LABORATORY TESTING BEEN PERFORMED?			
YES NO			
IF YES, PROVIDE RESULTS:			
Hemoglobin (gm/100ml): Date:			
Hematocrit: Date:			
Red blood cell (RBC) count: Date: White blood cell (WBC) count: Date:			
White blood cell differential count: Date:			
Platelet count: Date:			
9B. ARE THERE ANY OTHER SIGNIFICANT DIAGNOSTIC TEST FINDINGS AND/OR RESULTS? YES NO IF YES, PROVIDE TYPE OF TEST OR PROCEDURE, DATE AND RESULTS (brief summary): SECTION X - FUNCTIONAL IMPACT 0.00ES THE VETERAN'S HEMATOLOGIC OR LYMPHATIC CONDITION(S) IMPACT HIS OR HER ABILITY TO WORK? YES NO IF YES, DESCRIBE IMPACT OF EACH OF THE VETERAN'S HEMATOLOGIC AND/OR LYMPHATIC CONDITIONS, PROVIDING ONE OR MORE EXAMPLES:			
SECTION XI - REMARKS			
11. REMARKS (If any):			
SECTION XII - EXAMINER'S CERTIFICATION AND SIGNATURE			
CERTIFICATION - To the best of my knowledge, the information contained herein is accurate, complete and current.			
12A. Examiner's signature: 12B. Examiner's printed name and title (e.g. MD, DO, DDS, DMD, Ph.D, Psy.D, NP, PA-C):			
12C. Examiner's Area of Practice/Specialty (e.g. Cardiology, Orthopedics, Psychology/Psychiatry, General Practice): 12D. Date Signed:			
12E. Examiner's phone/fax numbers: 12F. National Provider Identifier (NPI) number: 12G. Medical license number and state:			
12F. National Provider Identifier (NPI) number: 12G. Medical license number and state:			
12H. Examiner's address:			