| Department of Veterans Affair | HEMATOLO | DISABILITY BENEFITS QUESTIONNAIRE | | | | |
|--|--|--|---|--|--|--|
| Name of Patient/Veteran | Patient/Veteran's Social | Security Number | Date of examination: | | | |
| IMPORTANT - THE DEPARTMENT OF VETERAN OF COMPLETING AND/OR SUBMITTING THIS FO | | PAY OR REIMBURSE AN | IY EXPENSES OR COST INCURRED IN THE PROCESS | | | |
| Note - The Veteran is applying to the U.S. Departm questionnaire as part of their evaluation in processis complete VA's review of the Veteran's application. \ questionnaire will be completed by the Veteran's | ng the Veteran's claim. VA m VA reserves the right to confi | ay obtain additional medica | al information, including an examination, if necessary, to | | | |
| Are you completing this Disability Benefits Question | nnaire at the request of: | | | | | |
| Veteran/Claimant | | | | | | |
| Third party (please list name(s) of organization | n(s) or individual(s)) | | | | | |
| Other: please describe | | | | | | |
| Are you a VA Healthcare provider? Yes | ○ No | | | | | |
| Is the Veteran regularly seen as a patient in your cli | inic? Yes | ○ No | | | | |
| Was the Veteran examined in person? Yes | ○ No | O | | | | |
| If no, how was the examination conducted? | | | | | | |
| in no, now was the examination conducted: | | | | | | |
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| | EVIDEN | ICE REVIEW | | | | |
| Evidence reviewed: | | | | | | |
| No records were reviewed | | | | | | |
| Records reviewed | | | | | | |
| Please identify the evidence reviewed (e.g. service | treatment records, VA treatm | nent records, private treatm | nent records) and the date range. | | | |
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| | SECTION | I - DIAGNOSIS | | | | |
| 1A. CHECK THE CLAIMED HEMATOLOGICAL AN | | ION(S) THAT DEDTAIN T | O THIS DRO: | | | |
| NOTE: These are the diagnoses determined during from a previous diagnosis for this condition, or if the comments section. Date of diagnosis can be the da | this current evaluation of the | claimed condition(s) listed cation due to the claimed c | d above. If there is no diagnosis, if the diagnosis is differen | | | |
| review or reported history. Agranulocytosis, acquired | | ICD code: | Date of diagnosis: | | | |
| | | | | | | |

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| Leukemia | | |
|--|---|--------------------------------|
| Chronic myelogenous leukemia (CML) (chronic myeloid leukemia or chronic granulocytic leukemia) | 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 | ICD code: | Date of diagnosis: |
| Active disease Treatment phase | | |
| Non-Hodgkin's lymphoma | ICD code: | Date of diagnosis: |
| Active disease Treatment phase Indolen | t and non-contiguous phase of low grade | NHL |
| Multiple myeloma | ICD code: | Date of diagnosis: |
| Monoclonal gammopathy of undetermined significance (MGUS) | 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 | ICD code: | Date of diagnosis: |
| Folic acid deficiency | ICD code: | Date of diagnosis: |
| Pernicious anemia or other Vitamin B12 deficiency anemia | ICD code: | Date of diagnosis: |
| Acquired hemolytic anemia | ICD code: | Date of diagnosis: |
| Other | ICD code: | Date of diagnosis: |
| AL amyloidosis (primary amyloidosis) | ICD code: | Date of diagnosis: |
| Immune thrombocytopenia | ICD code: | Date of diagnosis: |
| Polycythemia vera | ICD code: | Date of diagnosis: |
| Sickle cell anemia | ICD code: | Date of diagnosis: |
| Splenectomy | ICD code: | Date of diagnosis: |
| Are there complications such as systemic infections with encapsulated bacteria? | Yes No | |
| If Yes, complete SECTION VIII - OTHER PERTINENT PHYSICAL FINDINGS, CO | MPLICATIONS, CONDITIONS, SIGNS AN | ND/OR SYMPTOMS. |
| Injury to Spleen | ICD code: | Date of diagnosis: |
| If checked, complete SECTION VIII - OTHER PERTINENT PHYSICA | AL FINDINGS, COMPLICATIONS, COND | ITIONS, SIGNS AND/OR SYMPTOMS. |
| Adenitis, tuberculous (Also complete the Infectious Diseases (Other Than HIV-Related Illness, Chronic Fatigue Syndrome, or Tuberculosis) Disability Benefits Questionnaire). | ICD code: | 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: |

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| 1B. IF THERE ARE ADDITIONAL OR PRIOR DIAGNOSES THAT PERTAIN TO HEMATOLOGIC OR LYMPHATIC CONDITIONS, LIST USING ABOVE FORMAT: |
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| 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): |
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| 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 THROUBOCYTOPENIA CAUSED BY TREATMENT FOR A HEMATOLOGIC OR LYMPHATIC CONDITION, INCLUDING MEDICATION AND THE CONDITION THE MEDICATION IS USED TO TREAT: |
| WEDICATION AND THE CONDITION THE WEDICATION IS COLD TO TREAT. |
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| 2C. INDICATE THE STATUS OF THE PRIMARY HEMATOLOGIC OR LYMPHATIC CONDITION: |
| ACTIVE REMISSION NOT APPLICABLE |
| |
| 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 |
| Peripheral blood stem cell transplant Bone marrow stem cell transplant Other (specify) |
| |
| Other (specify) |

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| 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: |
| Date of most recent 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 LYMPHATIC |
| CONDITION? (Yes () No IF YES, COMPLETE THE FOLLOWING: |
| |
| 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? |
| O Yes O No IF YES, PROVIDE THE NAME OF THE OTHER HEMATOLOGIC OR LYMPHATIC CONDITION CAUSING THE SECONDARY ANEMIA: |
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| 10. ADI ADEIO ANELINA |
| 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 | | | | |
| 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 immune globulin |
| 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, MONOCLONAL GAMMOPATHY OF UNDETERMINED SIGNIFICANCE (MGUS), AGRANULOCYTOSIS, ACQUIRED, ESSENTIAL THROMBOCYTHEMIA, PRIMARY MYELOFIBROSIS, AND MYELODYSPLASTIC SYNDROMES |
| 5A. DOES THE VETERAN HAVE LEUKEMIA, MULTIPLE MYELOMA, MONOCLONAL GAMMOPATHY OF UNDETERMINED SIGNIFICANCE (MGUS), 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) |
| Monoclonal gammopathy of undetermined significance (MGUS) (complete 5C) |
| Agranulocytosis, acquired (complete 5D) |
| Essential thrombocythemia or primary myelofibrosis (complete 5E) |
| Myelodysplastic syndromes (complete 5F) |
| Other, specify |
| 5B. WHAT IS THE STATUS OF LEUKEMIA? |
| ○ ACTIVE ○ REMISSION |
| Asymptomatic, Rai Stage 0 |
| Requiring peripheral blood stem cell transplant |
| Requiring bone marrow stem cell transplant |
| Requiring continuous myelosuppressive therapy |
| |

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| Requiring continuous immunosuppressive therapy treatment | |
|---|---------|
| | |
| Requiring intermittent myelosuppressive therapy, or molecularly targeted therapy with tyrosine kinase inhibitors, or interferon treatment when not in agreemission | parent |
| In apparent remission on continuous molecularly targeted therapy with tyrosine kinase inhibitors | |
| 5C. WHAT IS THE STATUS OF MULTIPLE MYELOMA? | |
| Asymptomatic | |
| Monoclonal gammopathy of undetermined significance (MGUS) | |
| Smoldering multiple myeloma (SMM) | |
| Symptomatic (if checked, provide date of the diagnosis of symptomatic multiple myeloma) | |
| NOTE: Current validated biomarkers of symptomatic multiple myeloma, asymptomatic, smoldering or monoclonal gammopathy of undetermined significant are acceptable for the diagnosis of multiple myeloma as defined by the American Society of Hematology (ASH) and International Myeloma Working Group | |
| 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 (GN | И-CSF)) |
| Requiring continuous immunosuppressive therapy such as cyclosporine to maintain absolute neutrophil count (ANC) greater than 500/microliter (μl) b than 1000/μl | ut less |
| 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 every three months 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? | |
| GI. WINT TO THE CITATION OF INTELEST OF ENGINEER. | |
| Requiring peripheral blood stem cell transplant | |
| | |
| Requiring peripheral blood stem cell transplant | |
| Requiring peripheral blood stem cell transplant Requiring bone marrow stem cell transplant | |
| Requiring peripheral blood stem cell transplant Requiring bone marrow stem cell transplant Requiring chemotherapy | |

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| 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 |
| SECTION VI - POLYCYTHEMIA VERA |
| 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? |
| Yes No 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 |
| 8A. 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? |
| |
| If yes, describe (brief summary): Also if indicated, complete the appropriate questionnaire for each condition |
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| 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? | | | | |
|---|---|--|--|--|
| ○ Yes ○ No | | | | |
| IF YES, ALSO COMPLETE APPROPRIATE DERMATOLOGICAL I | DBQ | | | |
| SEC | CTION IX - DIAGNOSTIC TESTING | | | |
| NOTE: If testing has been performed and reflects Veteran's current count. | condition, no further testing is required. When appropriate, provide most recent complete blood | | | |
| 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): | | | |
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| SEC | CTION X - FUNCTIONAL IMPACT | | | |
| 10A. DOES THE VETERAN'S HEMATOLOGIC OR LYMPHATIC C | CONDITION(S) IMPACT HIS OR HER ABILITY TO WORK? | | | |
| Yes No | | | | |
| IF YES, DESCRIBE IMPACT OF EACH OF THE VETERAN'S HEM | MATOLOGIC AND/OR LYMPHATIC CONDITIONS, PROVIDING ONE OR MORE EXAMPLES: | | | |
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| SECTION XI - REMARKS | | | | |
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| 11A. REMARKS (If any): | | | | |
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| SECTIO | N XII - EXAN | MINER'S CERTIFICATION AND SIG | NATURE | |
| CERTIFICATION - To the best of my knowledge, the in | nformation cor | ntained herein is accurate, complete and cu | ırrent. | |
| PENALTY: The law provides severe penalties which in knowing it to be false, or for the fraudulent acceptance | clude fine or in of any payme | mprisonment, or both, for the willful submis nt to which you are not entitled. | sion of any sta | atement or evidence of a material fact, |
| 12A. Examiner's signature: 12. Examiner's printed name and title (e.g. MD, DO, DD | | DS, DMD, Ph.D, Psy.D, NP, PA-C): | | |
| 12C. Examiner's Area of Practice/Specialty (e.g. Cardi | ology, Orthope | edics, Psychology/Psychiatry, General Prac | ctice): | 12D. Date Signed: |
| 12E. Examiner's phone/fax numbers: | 12F. Nationa | al Provider Identifier (NPI) number: | 12G. Medica | I license number and state: |
| 12H. Examiner's address: | 1 | | - | |
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