



**DEPARTMENT OF VETERANS AFFAIRS  
Veterans Benefits Administration  
Regional Office**

**John James Rambo**

**VA File Number**

**000 00 0012**

**Represented By: VETERANS OF FOREIGN WARS OF THE US**

**Rating Decision**

**08/03/2020**

**INTRODUCTION**

The records reflect that you are a Veteran of the Vietnam War and Gulf War Era. You served in the Army from July 8, 1965 to July 31, 2020. You filed an original disability claim that was received on August 1, 2020. Based on a review of the evidence listed below, we have made the following decisions on your claim.

**DECISION**

1. Service connection for allergic rhinitis is denied.

**EVIDENCE**

- VA Form 21-526 EZ: Application for Disability Compensation and Related Compensation Benefits, February 11, 2020
- VA Form 21-0781, Statement In Support Of Claim For Service Connection for Post-Traumatic Stress Disorder (PTSD), received February 11, 2020
- VA Form 21-4138, Statement in Support of Claim, received February 11, 2020
- Service Treatment and Personnel Records received February 11, 2020, March 24, 2020, April 16, 2020, April 28, 2020 and June 3, 2020

- VA Contract Examinations for allergic rhinitis dated March 26, 2020.
- Section (§) 5103 Notice Response, received May 5, 2020
- DD214 and electronic review of service personnel file showing active duty service from July 8, 1965 to August 1, 2020

## **REASONS FOR DECISION**

### **1. Service connection for allergic rhinitis is denied.**

Service connection may be granted for a disability which began in military service or was caused by some event or experience in service. (38 CFR 3.303)

Service connection for allergic rhinitis is denied since this condition neither occurred in nor was caused by service. (38 CFR 3.303, 38 CFR 3.304)

Review of your service treatment records does not show a diagnosis of allergic rhinitis. Your claim of service connection for allergic rhinitis is denied.

### **REFERENCES:**

Title 38 of the Code of Federal Regulations, Pensions, Bonuses and Veterans' Relief contains the regulations of the Department of Veterans Affairs which govern entitlement to all veteran benefits. For additional information regarding applicable laws and regulations, please consult your local library, or visit us at our website, [www.va.gov](http://www.va.gov).



**SINUSITIS/RHINITIS AND OTHER CONDITIONS OF THE NOSE, THROAT,  
LARYNX AND PHARYNX DISABILITY BENEFITS QUESTIONNAIRE**

**IMPORTANT** - THE DEPARTMENT OF VETERANS AFFAIRS (VA) **WILL NOT PAY** OR **REIMBURSE** ANY EXPENSES OR COST INCURRED IN THE PROCESS OF COMPLETING AND/OR SUBMITTING THIS FORM. PLEASE READ THE PRIVACY ACT AND RESPONDENT BURDEN INFORMATION BEFORE COMPLETING FORM.

NAME OF PATIENT/VETERAN

John James Rambo

PATIENT/VETERAN'S SOCIAL SECURITY NUMBER

xxx-xx-0012

**NOTE TO PHYSICIAN** - Your patient is applying to the U.S. Department of Veterans Affairs (VA) for disability benefits. VA will consider the information you provide on this questionnaire as part of their evaluation in processing the Veteran's claim.

Is this questionnaire being completed in conjunction with a VA 21-2507, C&amp;P examination request?

☒ Yes ☐ No

How was the examination completed? Check all that apply:

☒ In-person examination☐ Records reviewed

Comments:

☒ Examination via approved telehealth☐ Other, please specify in comments box:**ACCEPTABLE CLINICAL EVIDENCE (ACE)**

INDICATE METHOD USED TO OBTAIN MEDICAL INFORMATION TO COMPLETE THIS DOCUMENT:

☐ Review of available records (without in-person or video telehealth examination) using the Acceptable Clinical Evidence (ACE) process because the existing medical evidence provided sufficient information on which to prepare the questionnaire and such an examination will likely provide no additional relevant evidence.☐ Review of available records in conjunction with an interview with the Veteran (without in-person or telehealth examination) using the ACE process because the existing medical evidence supplemented with an interview provided sufficient information on which to prepare the questionnaire and such an examination would likely provide no additional relevant evidence.**EVIDENCE REVIEW**

EVIDENCE REVIEWED (check all that apply):

☐ Not requested☐ No records were reviewed☐ VA claims file (hard copy paper C-file)☒ VA e-folder (VBMS or Virtual VA)☐ CPRS☐ Other (please identify other evidence reviewed):

EVIDENCE COMMENTS:

**SECTION I - DIAGNOSIS**

1A. DOES THE VETERAN NOW HAVE OR HAS HE OR SHE EVER BEEN DIAGNOSED WITH A SINUS, NOSE, THROAT, LARYNX OR PHARYNX CONDITION? *(This is the condition the Veteran is claiming or for which an exam has been requested.)*

☒ YES ☐ NO

1B. IF YES, SELECT THE VETERAN'S CONDITION *(check all that apply)*

<input type="checkbox"/> CHRONIC SINUSITIS	ICD Code: _____	Date of diagnosis: _____
<input checked="" type="checkbox"/> ALLERGIC RHINITIS	ICD Code: <u>J30.9</u>	Date of diagnosis: <u>01/01/2000</u>
<input type="checkbox"/> NON-ALLERGIC RHINITIS	ICD Code: _____	Date of diagnosis: _____
<input type="checkbox"/> BACTERIAL RHINITIS	ICD Code: _____	Date of diagnosis: _____
<input type="checkbox"/> GRANULOMATOUS RHINITIS	ICD Code: _____	Date of diagnosis: _____
<input type="checkbox"/> CHRONIC LARYNGITIS	ICD Code: _____	Date of diagnosis: _____
<input type="checkbox"/> LARYNGECTOMY	ICD Code: _____	Date of diagnosis: _____
<input type="checkbox"/> LARYNGEAL STENOSIS	ICD Code: _____	Date of diagnosis: _____
<input type="checkbox"/> APHONIA	ICD Code: _____	Date of diagnosis: _____
<input type="checkbox"/> PHARYNGEAL INJURY <i>(Describe):</i>	ICD Code: _____	Date of diagnosis: _____

<input type="checkbox"/> DEVIATED NASAL SEPTUM <i>(Traumatic)</i>	ICD Code: _____	Date of diagnosis: _____
<input type="checkbox"/> ANATOMICAL LOSS OF PART OF NOSE <i>(Complete Scars Benefits Questionnaire in lieu of this questionnaire)</i>	ICD Code: _____	Date of diagnosis: _____
<input type="checkbox"/> BENIGN OR MALIGNANT NEOPLASM OF SINUS, NOSE, THROAT, LARYNX OR PHARYNX	ICD Code: _____	Date of diagnosis: _____
<input checked="" type="checkbox"/> OTHER <i>(specify)</i>		
Other diagnosis #1 <u>Acute Sinusitis</u>	ICD Code: <u>J01.90</u>	Date of diagnosis: <u>01/01/2000</u>
Other diagnosis #2 _____	ICD Code: _____	Date of diagnosis: _____

1C. IF THERE ARE ADDITIONAL DIAGNOSES THAT PERTAIN TO THE SINUSES, NOSE, THROAT, LARYNX, OR PHARYNX CONDITION(S), LIST USING ABOVE FORMAT:

**SECTION II - MEDICAL HISTORY**

2. DESCRIBE THE HISTORY *(including onset and course)* OF THE VETERAN'S SINUS, NOSE, THROAT, LARYNX, OR PHARYNX CONDITION:

Onset date:

2000

Details of onset:

Veteran reports at onset, he had cold symptoms like congestion and runny nose. Veteran reports he was put on antibiotics, which did not help.

Course since onset:

Veteran reports he still gets allergic rhinitis symptoms. Veteran reports he has had sinus infections before as a result of untreated allergic rhinitis.

Current symptoms:

Veteran reports he gets allergic rhinitis symptoms of runny nose, cough and congestion at least once per month. Veteran reports having sniffles and congestion daily.

Current treatment:

Veteran reports taking claritin daily.

**SECTION III - NOSE, THROAT, LARYNX OR PHARYNX CONDITIONS**

3. DOES THE VETERAN HAVE ANY OF THE FOLLOWING NOSE, THROAT, LARYNX OR PHARYNX CONDITIONS?

☒ YES ☐ NO (If "No," proceed to Section IV) (If "Yes," check all that apply):

- ☒ Sinusitis (If checked, complete Part A below)
- ☒ Rhinitis (If checked, complete Part B below)
- ☐ Larynx or pharynx condition (If checked, complete Part C below)
- ☐ Deviated nasal septum (traumatic) (If checked, complete Part D below)
- ☐ Tumors or neoplasms (If checked, complete Part E below)
- ☐ Other nose, throat, larynx or pharynx conditions, pertinent physical findings or scars due to nose, throat, larynx or pharynx conditions. (If checked, complete Part F below)

**PART A - SINUSITIS**

A1. INDICATE THE SINUSES/TYPE OF SINUSITIS CURRENTLY AFFECTED BY THE VETERAN'S CHRONIC SINUSITIS (Check all that apply):

☐ NONE ☐ MAXILLARY ☐ FRONTAL ☐ ETHMOID ☐ SPHENOID ☐ PANSINUSITIS

A2. DOES THE VETERAN CURRENTLY HAVE ANY FINDINGS, SIGNS OR SYMPTOMS ATTRIBUTABLE TO CHRONIC SINUSITIS?

☐ YES ☒ NO

(If "Yes," check all that apply)

- ☐ Chronic sinusitis detected only by imaging studies (See Diagnostic Testing Section)
- ☐ Episodes of sinusitis
- ☐ Near constant sinusitis (If checked, describe frequency): \_\_\_\_\_
- ☐ Headaches
- ☐ Pain of affected sinus
- ☐ Tenderness of affected sinus
- ☐ Purulent discharge
- ☐ Crusting
- ☐ Other (describe): \_\_\_\_\_

FOR ALL CHECKED CONDITIONS, DESCRIBE: \_\_\_\_\_

A3. HAS THE VETERAN HAD **NON-INCAPACITATING** EPISODES OF SINUSITIS CHARACTERIZED BY HEADACHES, PAIN AND PURULENT DISCHARGE OR CRUSTING IN THE PAST 12 MONTHS?☐ YES ☒ NO

(If "Yes," provide the total number of non-incapacitating episodes over the past 12 months):

☐ 1 ☐ 2 ☐ 3 ☐ 4 ☐ 5 ☐ 6 ☐ 7 or moreA4. HAS THE VETERAN HAD **INCAPACITATING** EPISODES OF SINUSITIS REQUIRING PROLONGED (4 to 6 weeks) OF ANTIBIOTICS TREATMENT IN THE PAST 12 MONTHS?**NOTE** - For VA purposes, an incapacitating episode of sinusitis means one that requires bed rest and treatment prescribed by a physician.☐ YES ☒ NO

(If "Yes," provide the total number of incapacitating episodes of sinusitis requiring prolonged (4 to 6 weeks) of antibiotic treatment over the past 12 months):

☐ 1 ☐ 2 ☐ 3 or more

A5. HAS THE VETERAN HAD SINUS SURGERY?

☐ YES ☒ NO

(If "Yes," specify type of surgery):

- ☐ Radical (open sinus surgery) ☐ Endoscopic ☐ Other: \_\_\_\_\_
- (Type of procedure, sinuses operated on and side(s)): \_\_\_\_\_
- (Date(s) of surgery (if repeated sinus surgery, provide all dates of surgery)): \_\_\_\_\_

A6. IF VETERAN HAS HAD RADICAL SINUS SURGERY, DID CHRONIC OSTEOMYELITIS FOLLOW THE SURGERY?

☐ YES ☐ NO (If "Yes," complete Osteomyelitis Questionnaire)

A7. HAS THE VETERAN HAD REPEATED SINUS-RELATED SURGICAL PROCEDURES PERFORMED?

☐ YES ☒ NO**PART B - RHINITIS**

B1. IS THERE GREATER THAN 50% OBSTRUCTION OF THE NASAL PASSAGE ON BOTH SIDES DUE TO RHINITIS?

☒ YES ☐ NO

B2. IS THERE COMPLETE OBSTRUCTION ON THE LEFT SIDE DUE TO RHINITIS?

☐ YES ☒ NO

B3. IS THERE COMPLETE OBSTRUCTION ON THE RIGHT SIDE DUE TO RHINITIS?

☐ YES ☒ NO

**SECTION III - NOSE, THROAT, LARYNX OR PHARYNX CONDITIONS (Continued)****PART B - RHINITIS (Continued)**

B4. IS THERE PERMANENT HYPERTROPHY OF THE NASAL TURBINATES?

☐ YES ☒ NO

B5. ARE THERE NASAL POLYPS?

☐ YES ☒ NO

B6. DOES THE VETERAN HAVE ANY OF THE FOLLOWING GRANULOMATOUS CONDITIONS?

☐ YES ☒ NO (If "Yes," check all that apply)☐ Granulomatous rhinitis ☐ Rhinoscleroma ☐ Wegener's granulomatosis ☐ Lethal midline granuloma☐ Other granulomatous infection (Describe): \_\_\_\_\_**PART C - LARYNX AND PHARYNX CONDITIONS**

C1. DOES THE VETERAN HAVE CHRONIC LARYNGITIS?

☐ YES ☐ NO

(If "Yes," does the Veteran have any of the following symptoms due to chronic laryngitis?)

☐ YES ☐ NO (If "Yes," check all that apply)☐ Hoarseness (If checked, describe frequency): \_\_\_\_\_☐ Inflammation of vocal cords☐ Inflammation of mucous membrane☐ Thickening of vocal chords☐ Nodules of vocal chords☐ Submucous infiltration of vocal chords☐ Vocal chord polyps☐ Other (describe): \_\_\_\_\_

C2. HAS THE VETERAN HAD A LARYNGECTOMY?

☐ YES ☐ NO (If "Yes," specify)☐ Total laryngectomy☐ Partial laryngectomy

(If checked, does the Veteran have any residuals of the partial laryngectomy?)

☐ YES ☐ NO

(If "Yes," describe): \_\_\_\_\_

C3. DOES THE VETERAN HAVE LARYNGEAL STENOSIS, INCLUDING RESIDUALS OF LARYNGEAL TRAUMA (unilateral or bilateral)?

☐ YES ☐ NO (If "Yes," assess for upper airway obstruction with pulmonary function testing to include Flow-Volume Loop, and provide results in Diagnostic Testing Section)

C4. DOES THE VETERAN HAVE COMPLETE ORGANIC APHONIA?

☐ YES ☐ NO (If "Yes," check all that apply)☐ Constant inability to speak above a whisper☐ Constant inability to communicate by speech☐ Other (describe): \_\_\_\_\_

C5. DOES THE VETERAN HAVE INCOMPLETE ORGANIC APHONIA?

☐ YES ☐ NO (If "Yes," check all that apply)☐ Hoarseness (If checked, describe frequency): \_\_\_\_\_☐ Inflammation of vocal cords☐ Inflammation of mucous membrane☐ Thickening of vocal chords☐ Nodules of vocal chords☐ Submucous infiltration of vocal chords☐ Vocal chord polyps☐ Other (describe): \_\_\_\_\_

C6. HAS THE VETERAN HAD A PERMANENT TRACHEOSTOMY?

☐ YES ☐ NO (If "Yes," describe reason for tracheostomy and potential for decannulation):

**SECTION III - NOSE, THROAT, LARYNX OR PHARYNX CONDITIONS (Continued)****PART C - LARYNX AND PHARYNX CONDITIONS**

C7. HAS THE VETERAN HAD AN INJURY TO THE PHARYNX?

☐ YES ☐ NO (If "Yes," check all findings, signs and symptoms that apply):

- ☐ Obstruction of the pharynx
- ☐ Obstruction of the nasopharynx
- ☐ Stricture of the pharynx
- ☐ Stricture of the nasopharynx
- ☐ Absence of the soft palate secondary to trauma
- ☐ Absence of the soft palate secondary to chemical burn
- ☐ Absence of the soft palate secondary to granulomatous disease
- ☐ Paralysis of the soft palate
- ☐ Swallowing difficulty
- ☐ Nasal regurgitation
- ☐ Speech impairment
- ☐ Other (describe): \_\_\_\_\_

C8. DOES THE VETERAN HAVE VOCAL CHORD PARALYSIS OR ANY OTHER PHARYNGEAL OR LARYNGEAL CONDITIONS?

☐ YES ☐ NO (If "Yes," describe): \_\_\_\_\_**PART D - DEVIATED NASAL SEPTUM (TRAUMATIC)**

D1. IS THERE AT LEAST 50% OBSTRUCTION OF THE NASAL PASSAGE ON BOTH SIDES DUE TO TRAUMATIC SEPTAL DEVIATION?

☐ YES ☐ NO

D2. IS THE VETERAN'S DEVIATED SEPTUM TRAUMATIC?

☐ YES ☐ NO

D3. IS THERE COMPLETE OBSTRUCTION ON LEFT SIDE DUE TO TRAUMATIC SEPTAL DEVIATION?

☐ YES ☐ NO

D4. IS THERE COMPLETE OBSTRUCTION ON RIGHT SIDE DUE TO TRAUMATIC SEPTAL DEVIATION?

☐ YES ☐ NO**PART E - TUMORS AND NEOPLASMS**

E1. DOES THE VETERAN HAVE A BENIGN OR MALIGNANT NEOPLASM OR METASTASES RELATED TO ANY OF THE DIAGNOSES IN THE DIAGNOSIS SECTION?

☐ YES ☐ NO (If "Yes," complete the following section)

E2. IS THE NEOPLASM:

☐ BENIGN ☐ MALIGNANT

E3. HAS THE VETERAN COMPLETED TREATMENT OR IS THE VETERAN CURRENTLY UNDERGOING TREATMENT FOR A BENIGN OR MALIGNANT NEOPLASM OR METASTASES?

☐ 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☐ 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): \_\_\_\_\_

E4. DOES THE VETERAN CURRENTLY HAVE ANY RESIDUAL CONDITIONS OR COMPLICATIONS DUE TO THE NEOPLASM (including metastases) OR ITS TREATMENT, OTHER THAN THOSE ALREADY DOCUMENTED IN THE REPORT ABOVE?

☐ YES ☐ NO (If "Yes," list residual conditions and complications (brief summary)):



**SECTION III - NOSE, THROAT, LARYNX OR PHARYNX CONDITIONS (Continued)****PART E - TUMORS AND NEOPLASMS (Continued)**

E5. IF THERE ARE ADDITIONAL BENIGN OR MALIGNANT NEOPLASMS OR METASTASES RELATED TO ANY OF THE DIAGNOSES IN THE DIAGNOSIS SECTION, DESCRIBE USING THE ABOVE FORMAT:

**PART F - OTHER PERTINENT PHYSICAL FINDINGS, COMPLICATIONS, CONDITIONS, SIGNS, SYMPTOMS, AND SCARS**

F1. DOES THE VETERAN HAVE ANY OTHER PERTINENT PHYSICAL FINDINGS, COMPLICATIONS, CONDITIONS, SIGNS OR SYMPTOMS RELATED TO THE CONDITIONS LISTED IN THE DIAGNOSIS SECTION ABOVE?

☐ YES ☐ NO

IF YES, DESCRIBE (*brief summary*):

F2. DOES THE VETERAN HAVE ANY SCARS (*surgical or otherwise*) RELATED TO ANY CONDITIONS OR TO THE TREATMENT OF ANY CONDITIONS LISTED IN THE DIAGNOSIS SECTION ABOVE?

☐ YES ☐ NO

IF YES, ARE ANY OF THESE SCARS PAINFUL OR UNSTABLE; HAVE A TOTAL AREA EQUAL TO OR GREATER THAN 39 SQUARE CM (*6 square inches*); OR ARE LOCATED ON THE HEAD, FACE OR NECK? (An "unstable scar" is one where, for any reason, there is frequent loss of covering of the skin over the scar.)

☐ YES ☐ NO

IF YES, ALSO COMPLETE VA FORM 21-0960F-1, SCARS/DISFIGUREMENT.

IF NO, PROVIDE LOCATION AND MEASUREMENTS OF SCAR IN CENTIMETERS.

LOCATION: \_\_\_\_\_ MEASUREMENTS: length \_\_\_\_\_ cm X width \_\_\_\_\_ cm.

**NOTE:** If there are multiple scars, enter additional locations and measurements in Comment section below. It is not necessary to also complete a Scars DBQ.

F3. COMMENTS, IF ANY:

F4. DOES THE VETERAN HAVE LOSS OF PART OF THE NOSE OR OTHER SCARS OF THE NOSE EXPOSING BOTH NASAL PASSAGES?

☐ YES ☐ NO

F5. DOES THE VETERAN HAVE LOSS OF PART OF THE NOSE OR OTHER SCARS CAUSING LOSS OF PART OF ONE ALA?

☐ YES ☐ NO

F6. DOES THE VETERAN HAVE LOSS OF PART OF THE NOSE OR OTHER SCARS CAUSING ANY OTHER DISFIGUREMENT?

☐ YES ☐ NO

**SECTION IV - DIAGNOSTIC TESTING**

**NOTE** - If testing has been performed and reflects the Veteran's current condition, repeat testing is not required. Specific diagnostic testing is not required for many conditions, but if performed, record in this section.

**4A. HAVE IMAGING STUDIES OF THE SINUSES OR OTHER AREAS BEEN PERFORMED?**

☒ YES ☐ NO

*(If "Yes," check all that apply)*

☐ Magnetic resonance imaging (MRI) Date: \_\_\_\_\_ Results: \_\_\_\_\_

☐ Computed tomography (CT) Date: \_\_\_\_\_ Results: \_\_\_\_\_

☒ X-rays: x-ray paranasal sinuses Date: 08/17/2020 Results: normal sinus cavities

☐ Other: \_\_\_\_\_ Date: \_\_\_\_\_ Results: \_\_\_\_\_

**4B. HAS ENDOSCOPY BEEN PERFORMED?**

☐ YES ☒ NO

*(If "Yes," check all that apply):*

☐ Nasal endoscopy Date: \_\_\_\_\_ Results: \_\_\_\_\_

☐ Laryngeal endoscopy Date: \_\_\_\_\_ Results: \_\_\_\_\_

☐ Bronchoscopy Date: \_\_\_\_\_ Results: \_\_\_\_\_

☐ Other endoscopy Date: \_\_\_\_\_ Results: \_\_\_\_\_

**4C. HAS THE VETERAN HAD A BIOPSY OF THE LARYNX OR PHARYNX?**

☐ YES ☒ NO

*(If "Yes," complete the following):*

Site of biopsy: \_\_\_\_\_ Date: \_\_\_\_\_

Results: ☐ Benign ☐ Pre-malignant ☐ Malignant

Describe results: \_\_\_\_\_

**4D. HAS THE VETERAN HAD PULMONARY FUNCTION TESTING TO ASSESS FOR UPPER AIRWAY OBSTRUCTION DUE TO LARYNGEAL STENOSIS?**

☐ YES ☒ NO

*If "Yes," indicate results:*

☐ FEV-1 of 71 to 80% predicted

☐ FEV-1 of 56 to 70% predicted

☐ FEV-1 of 40 to 55% predicted

☐ FEV-1 less than 40% predicted

*Is the Flow-Volume Loop compatible with upper airway obstruction?*

☐ YES ☐ NO

**4E. 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 V - FUNCTIONAL IMPACT AND REMARKS**

5A. DOES THE VETERAN'S SINUS, NOSE, THROAT, LARYNX OR PHARYNX CONDITION IMPACT HIS OR HER ABILITY TO WORK?

☒ YES ☐ NO (If "Yes," describe impact of each of the veteran's sinus, nose, throat, larynx or pharynx conditions, providing one or more examples):

Current OR if retired/unemployed, previous occupation:

unemployed, previously army

0-1 week work time lost in last 12 months

Veteran reports it is harder to run, due to chronic congestion and sniffles.

5B. REMARKS (If any)

Is there a need for the Veteran/Service Member to follow up with his or her primary care provider regarding any findings in this examination (not limited to claimed condition(s))? No

Is the Veteran homeless? No

Veteran was instructed to send all personal medical records to the VA Evidence Intake Center if applicable, for proper submission into VBMS.

NOTE: VA may request additional medical information, including additional examinations if necessary to complete VA's review of the application.

**SECTION VII - PHYSICIAN'S CERTIFICATION AND SIGNATURE****CERTIFICATION** - To the best of my knowledge, the information contained herein is accurate, complete and current.

7A. PHYSICIAN'S SIGNATURE

John Smith

7B. PHYSICIAN'S PRINTED NAME

John Smith

7C. DATE SIGNED

03/26/2020

7D. PHYSICIAN'S PHONE AND FAX NUMBERS

619-400-5555

7E. NATIONAL PROVIDER IDENTIFIER (NPI) NUMBER

123456

7F. PHYSICIAN'S ADDRESS

8810 Rio San Diego Dr

NOTE - VA may request additional medical information, including additional examinations, if necessary to complete VA's review of the veteran's application.

**PRIVACY ACT NOTICE:** VA will not disclose information collected on this form to any source other than what has been authorized under the Privacy Act of 1974 or Title 38, Code of Federal Regulations 1.576 for routine uses (i.e., civil or criminal law enforcement, congressional communications, epidemiological or research studies, the collection of money owed to the United States, litigation in which the United States is a party or has an interest, the administration of VA programs and delivery of VA benefits, verification of identity and status, and personnel administration) as identified in the VA system of records, 58/VA21/22/28, Compensation, Pension, Education and Vocational Rehabilitation and Employment Records - VA, published in the Federal Register. Your obligation to respond is voluntary. VA uses your SSN to identify your claim file. Providing your SSN will help ensure that your records are properly associated with your claim file. Giving us your SSN account information is voluntary. Refusal to provide your SSN by itself will not result in the denial of benefits. VA will not deny an individual benefits for refusing to provide his or her SSN unless the disclosure of the SSN is required by a Federal Statute of law in effect prior to January 1, 1975, and still in effect. The requested information is considered relevant and necessary to determine maximum benefits under the law. The responses you submit are considered confidential (38 U.S.C. 5701). Information submitted is subject to verification through computer matching programs with other agencies.

**RESPONDENT BURDEN:** We need this information to determine entitlement to benefits (38 U.S.C. 501). Title 38, United States Code, allows us to ask for this information. We estimate that you will need an average of 15 minutes to review the instructions, find the information, and complete the form. VA cannot conduct or sponsor a collection of information unless a valid OMB control number is displayed. You are not required to respond to a collection of information if this number is not displayed. Valid OMB control numbers can be located on the OMB Internet Page at [www.reginfo.gov/public/do/PRAMain](http://www.reginfo.gov/public/do/PRAMain). If desired, you can call 1-800-827-1000 to get information on where to send comments or suggestions about this form.



**INTERNAL VETERANS AFFAIRS USE  
SHOULDER AND ARM CONDITIONS  
DISABILITY BENEFITS QUESTIONNAIRE**

Name of Claimant/Veteran:

John James Rambo

Claimant/Veteran's Social Security Number:

xxxxx0012

Date of Examination:

08/26/2020

Note to examiner - The Veteran is applying to the U.S. Department of Veterans Affairs (VA) for disability benefits. VA will consider the information you provide on this questionnaire as part of their evaluation in processing the Veteran's claim.

Is this questionnaire being completed in conjunction with VA 21-2507, C&amp;P examination request?

☒ Yes ☐ No

How was the examination completed? (check all that apply)

☒ In-person examination☒ Records reviewed☐ Examination via approved video telehealth

Comments:

☐ Other, please specify in comments box:**ACCEPTABLE CLINICAL EVIDENCE (ACE)**

Indicate the method used to obtain medical information to complete this document:

☒ Review of available records (without in-person or video telehealth examination) using the Acceptable Clinical Evidence (ACE) process because the existing medical evidence provided sufficient information on which to prepare the questionnaire and such an examination will likely provide no additional relevant evidence.☐ Review of available records in conjunction with an interview with the Veteran (without in-person or telehealth examination) using the ACE process because the existing medical evidence supplemented with an interview provided sufficient information on which to prepare the questionnaire and such an examination would likely provide no additional relevant evidence.**EVIDENCE REVIEW**

Evidence reviewed (check all that apply):

☐ Not requested☐ VA electronic health record☐ VA claims file (hard copy paper C-file)☐ No records were reviewed☒ VA e-folder☒ Other (please identify other evidence reviewed):

Evidence comments:

JLV

**DOMINANT HAND**

Dominant hand:

☒ Right☐ Left☐ Ambidextrous**SECTION I - DIAGNOSIS**

Note: These are condition(s) for which an evaluation has been requested on the exam request form (Internal VA) or for which the Veteran has requested medical evidence be provided for submission to VA.

1A. List the claimed conditions that pertain to this questionnaire: Right shoulder injury with degenerative changes

Note: These are the diagnoses determined during this current evaluation of the claimed condition(s) listed above. If there is no diagnosis, if the diagnosis is different from a previous diagnosis for this condition, or if there is a diagnosis of a complication due to the claimed condition, explain your findings and reasons in the remarks section. Date of diagnosis can be the date of the evaluation if the clinician is making the initial diagnosis or an approximate date determined through record review or reported history.

1B. Select diagnoses associated with the claimed condition(s) (check all that apply):

☐ The Veteran does not have a current diagnosis associated with any claimed conditions listed above. (Explain your findings and reasons in the remarks section)

	Side affected:			ICD Code:	Date of diagnosis:	
	Right	Left	Both		Right:	Left:
<input type="checkbox"/> Shoulder strain	<input type="checkbox"/> Right	<input type="checkbox"/> Left	<input type="checkbox"/> Both		Right: _____	Left: _____
<input type="checkbox"/> Shoulder impingement syndrome	<input type="checkbox"/> Right	<input type="checkbox"/> Left	<input type="checkbox"/> Both		Right: _____	Left: _____
<input type="checkbox"/> Bicipital tendonitis	<input type="checkbox"/> Right	<input type="checkbox"/> Left	<input type="checkbox"/> Both		Right: _____	Left: _____
<input type="checkbox"/> Bicipital tendon tear	<input type="checkbox"/> Right	<input type="checkbox"/> Left	<input type="checkbox"/> Both		Right: _____	Left: _____
<input checked="" type="checkbox"/> Rotator cuff tendonitis	<input checked="" type="checkbox"/> Right	<input type="checkbox"/> Left	<input type="checkbox"/> Both		Right: s46.092s 200	Left: _____

## SECTION I - DIAGNOSIS (continued)

	Side affected:			ICD Code:	Date of diagnosis:		
	Right	Left	Both		Right:	Left:	
<input type="checkbox"/> Rotator cuff tear	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<div></div>			
<input type="checkbox"/> Labral tear, including SLAP (superior labral anterior-posterior lesion)	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<div></div>			
<input type="checkbox"/> Subacromial/subdeltoid bursitis	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<div></div>			
<input type="checkbox"/> Glenohumeral joint osteoarthritis	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<div></div>			
<input type="checkbox"/> Acromioclavicular joint osteoarthritis	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<div></div>			
<input type="checkbox"/> Ankylosis of glenohumeral articulations (shoulder joint)	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<div></div>			
<input type="checkbox"/> Glenohumeral joint instability	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<div></div>			
<input type="checkbox"/> Glenohumeral joint dislocation/recurrent dislocation	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<div></div>			
<input type="checkbox"/> Shoulder joint replacement (total shoulder arthroplasty/hemiarthroplasty)	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<div></div>			
<input type="checkbox"/> Acromioclavicular joint separation	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<div></div>			
<input type="checkbox"/> Degenerative arthritis, other than post-traumatic	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<div></div>			
<input type="checkbox"/> Arthritis, gonorrheal	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<div></div>			
<input type="checkbox"/> Arthritis, pneumococcic	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<div></div>			
<input type="checkbox"/> Arthritis, streptococcic	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<div></div>			
<input type="checkbox"/> Arthritis, syphilitic	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<div></div>			
<input type="checkbox"/> Arthritis, rheumatoid (multi-joints)	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<div></div>			
<input type="checkbox"/> Post-traumatic arthritis	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<div></div>			
<input type="checkbox"/> Arthritis, typhoid	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<div></div>			
<input type="checkbox"/> Other specified forms of arthropathy (excluding gout) (specify)	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<div></div>			
<hr/>							
<input type="checkbox"/> Osteoporosis, residuals of	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<div></div>			
<input type="checkbox"/> Osteomalacia, residuals of	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<div></div>			
<input type="checkbox"/> Bones, neoplasm, benign	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<div></div>			
<input type="checkbox"/> Osteitis deformans	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<div></div>			
<input type="checkbox"/> Gout	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<div></div>			
<input type="checkbox"/> Bursitis	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<div></div>			
<input type="checkbox"/> Myositis	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<div></div>			
<input type="checkbox"/> Heterotopic ossification	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<div></div>			
<input type="checkbox"/> Tendinopathy (select one if known)	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<div></div>			
<input type="checkbox"/> Tendinitis	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<div></div>			
<input type="checkbox"/> Tendinosis	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<div></div>			
<input type="checkbox"/> Tenosynovitis	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<div></div>			
<input type="checkbox"/> Inflammatory - other types (specify)	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<div></div>			
<hr/>							
<input type="checkbox"/> Other (specify)	<div></div>						
<div>Other diagnosis #1 <div></div></div>							
Side affected:	<input type="checkbox"/> Right	<input type="checkbox"/> Left	<input type="checkbox"/> Both	ICD Code: <div></div>	Date of diagnosis:	Right: <div></div>	Left: <div></div>
<div>Other diagnosis #2 <div></div></div>							
Side affected:	<input type="checkbox"/> Right	<input type="checkbox"/> Left	<input type="checkbox"/> Both	ICD Code: <div></div>	Date of diagnosis:	Right: <div></div>	Left: <div></div>
If there are additional diagnoses that pertain to shoulder and/or arm conditions, list using above format:							
<div></div>							

## SECTION II - MEDICAL HISTORY

2A. Describe the history (including onset and course) of the Veteran's shoulder and/or arm condition (brief summary):

History of R shoulder pain since 2013. Presents with limitations in ROM, strength, posture, function and endurance. He reports occasional pain and stiffness.

**SECTION II - MEDICAL HISTORY (continued)**

2B. Does the Veteran report flare-ups of the shoulder and/or arm? ☒ Yes ☐ No

If yes, document the Veteran's description of the flare-ups he or she experiences, including the frequency, duration, characteristics, precipitating and alleviating factors, severity and/or extent of functional impairment he or she experiences during a flare-up of symptoms:

Patient reports pain with movement.

2C. Does the Veteran report having any functional loss or functional impairment of the joint or extremity being evaluated on this questionnaire, including but not limited to after repeated use over time? ☐ Yes ☒ No

If yes, document the Veteran's description of functional loss or functional impairment in his/her own words:

**SECTION III - RANGE OF MOTION (ROM) AND FUNCTIONAL LIMITATION**

There are several separate parameters requested for describing function of a joint. The question "Does this ROM contribute to a functional loss?" asks if there is a functional loss that can be ascribed to any documented loss of range of motion; and, unlike later questions, does not take into account the numerous other factors to be considered. Subsequent questions take into account additional factors such as pain, fatigue, weakness, lack of endurance, or incoordination. If there is pain noted on examination, it is important to understand whether or not that pain itself contributes to functional loss. Ideally, a claimant would be seen immediately after repetitive use over time or during a flare-up; however, this is not always feasible.

Information regarding joint function on repetitive use is broken up into two subsets. The first subset is based on observed repetitive use, and the second is based on functional loss associated with repeated use over time. The observed repetitive use section initially asks for objective findings after three or more repetitions of range of motion testing. The second subset provides a more global picture of functional loss associated with repetitive use over time. The latter takes into account medical probability of additional functional loss as a global view. This takes into account not only the objective findings noted on the examination, but also the subjective history provided by the claimant, as well as review of the available medical evidence.

Optimally, a description of any additional loss of function should be provided - such as what the degrees of range of motion would be opined to look like after repetitive use over time. However, when this is not feasible, an "as clear as possible" description of that loss should be provided. This same information (minus the three repetitions) is asked to be provided with regards to flare-ups.

**Right shoulder****Left shoulder****3A. Initial ROM measurements****3A. Initial ROM measurements**

☐ All normal ☒ Abnormal or outside of normal range  
☐ Unable to test ☐ Not indicated

☒ All normal ☐ Abnormal or outside of normal range  
☐ Unable to test ☐ Not indicated

If "Unable to test" or "Not indicated" please explain:

If "Unable to test" or "Not indicated" please explain:

If ROM is outside of "normal" range, but is normal for the Veteran (for reason other than a hip/thigh condition, such as age, body habitus, neurologic disease), please describe:

If ROM is outside of "normal" range, but is normal for the Veteran (for reason other than a hip/thigh condition, such as age, body habitus, neurologic disease), please describe:

If abnormal, does the range of motion itself contribute to a functional loss? (if yes, please explain) ☐ Yes ☒ No

If abnormal, does the range of motion itself contribute to a functional loss? (if yes, please explain) ☐ Yes ☐ No

Note: For any joint condition, examiners should address pain on both passive and active motion, and on both weight-bearing and nonweight-bearing. Examiners should also test the contralateral joint (unless medically contraindicated). If testing cannot be performed or is medically contraindicated (such as it may cause the Veteran severe pain or the risk of further injury), an explanation must be given below. Please note any characteristics of pain observed on examination (such as facial expression or wincing on pressure or manipulation).

Can testing be performed? ☒ Yes ☐ No If no, provide an explanation:

Can testing be performed? ☒ Yes ☐ No If no, provide an explanation:

If this is the unclaimed joint, is it: ☐ Damaged ☒ Undamaged

If this is the unclaimed joint, is it: ☐ Damaged ☒ Undamaged

If undamaged, range of motion testing must be conducted.

If undamaged, range of motion testing must be conducted.

### SECTION III - RANGE OF MOTION (ROM) AND FUNCTIONAL LIMITATION (continued)

3A. Initial ROM measurements (continued)	3A. Initial ROM measurements (continued)
<p>Active Range of Motion (ROM) - Perform active range of motion and provide the ROM values.</p> <p>Flexion endpoint (180 degrees):                      0-140                      degrees</p> <p>Abduction endpoint (180 degrees):                      0-130                      degrees</p> <p>Internal rotation endpoint (90 degrees):                      0-65                      degrees</p> <p>External rotation endpoint (90 degrees):                      0-70                      degrees</p> <p>If noted on examination, which ROM exhibited pain? (select all that apply):</p> <p><input type="checkbox"/> Flexion                      <input type="checkbox"/> Internal rotation</p> <p><input type="checkbox"/> Abduction                      <input type="checkbox"/> External rotation</p> <p>If any limitation of motion is specifically attributable to pain, weakness, fatigability, incoordination, or other; please note the degree(s) in which limitation of motion is specifically attributable to the factors identified and describe.</p> <p>_____ Flexion degree endpoint (if different than above)</p> <p>_____ Abduction degree endpoint (if different than above)</p> <p>_____ Internal rotation degree endpoint (if different than above)</p> <p>_____ External rotation degree endpoint (if different than above)</p>	<p>Active Range of Motion (ROM) - Perform active range of motion and provide the ROM values.</p> <p>Flexion endpoint (180 degrees):                      0-180                      degrees</p> <p>Abduction endpoint (180 degrees):                      0-180                      degrees</p> <p>Internal rotation endpoint (90 degrees):                      0-90                      degrees</p> <p>External rotation endpoint (90 degrees):                      0-90                      degrees</p> <p>If noted on examination, which ROM exhibited pain? (select all that apply):</p> <p><input type="checkbox"/> Flexion                      <input type="checkbox"/> Internal rotation</p> <p><input type="checkbox"/> Abduction                      <input type="checkbox"/> External rotation</p> <p>If any limitation of motion is specifically attributable to pain, weakness, fatigability, incoordination, or other; please note the degree(s) in which limitation of motion is specifically attributable to the factors identified and describe.</p> <p>_____ Flexion degree endpoint (if different than above)</p> <p>_____ Abduction degree endpoint (if different than above)</p> <p>_____ Internal rotation degree endpoint (if different than above)</p> <p>_____ External rotation degree endpoint (if different than above)</p>
<p>Passive Range of Motion - Perform passive ROM and provide the ROM values.</p> <p>Flexion endpoint (180 degrees):                      _____ degrees    <input type="checkbox"/> Same as active ROM</p> <p>Abduction endpoint (180 degrees):                      _____ degrees    <input type="checkbox"/> Same as active ROM</p> <p>Internal rotation endpoint (90 degrees):                      _____ degrees    <input type="checkbox"/> Same as active ROM</p> <p>External rotation endpoint (90 degrees):                      _____ degrees    <input type="checkbox"/> Same as active ROM</p> <p>If noted on examination, which ROM exhibited pain? (select all that apply):</p> <p><input type="checkbox"/> Flexion                      <input type="checkbox"/> Internal rotation</p> <p><input type="checkbox"/> Abduction                      <input type="checkbox"/> External rotation</p> <p>If any limitation of motion is specifically attributable to pain, weakness, fatigability, incoordination, or other; please note the degree(s) in which limitation of motion is specifically attributable to the factors identified and describe.</p> <p>_____ Flexion degree endpoint (if different than above)</p> <p>_____ Abduction degree endpoint (if different than above)</p> <p>_____ Internal rotation degree endpoint (if different than above)</p> <p>_____ External rotation degree endpoint (if different than above)</p>	<p>Passive Range of Motion - Perform passive ROM and provide the ROM values.</p> <p>Flexion endpoint (180 degrees):                      _____ degrees    <input type="checkbox"/> Same as active ROM</p> <p>Abduction endpoint (180 degrees):                      _____ degrees    <input type="checkbox"/> Same as active ROM</p> <p>Internal rotation endpoint (90 degrees):                      _____ degrees    <input type="checkbox"/> Same as active ROM</p> <p>External rotation endpoint (90 degrees):                      _____ degrees    <input type="checkbox"/> Same as active ROM</p> <p>If noted on examination, which ROM exhibited pain? (select all that apply):</p> <p><input type="checkbox"/> Flexion                      <input type="checkbox"/> Internal rotation</p> <p><input type="checkbox"/> Abduction                      <input type="checkbox"/> External rotation</p> <p>If any limitation of motion is specifically attributable to pain, weakness, fatigability, incoordination, or other; please note the degree(s) in which limitation of motion is specifically attributable to the factors identified and describe.</p> <p>_____ Flexion degree endpoint (if different than above)</p> <p>_____ Abduction degree endpoint (if different than above)</p> <p>_____ Internal rotation degree endpoint (if different than above)</p> <p>_____ External rotation degree endpoint (if different than above)</p>
<p>Is there evidence of pain?    <input checked="" type="checkbox"/> Yes    <input type="checkbox"/> No    If yes check all that apply.</p> <p><input type="checkbox"/> Weight-bearing                      <input type="checkbox"/> Nonweight-bearing</p> <p><input checked="" type="checkbox"/> Active motion                      <input type="checkbox"/> Passive motion</p> <p><input type="checkbox"/> On rest/non-movement    <input type="checkbox"/> Does not result in/cause functional loss</p> <p><input type="checkbox"/> Causes functional loss (if checked describe in the comments box below)</p>	<p>Is there evidence of pain?    <input type="checkbox"/> Yes    <input checked="" type="checkbox"/> No    If yes check all that apply.</p> <p><input type="checkbox"/> Weight-bearing                      <input type="checkbox"/> Nonweight-bearing</p> <p><input type="checkbox"/> Active motion                      <input type="checkbox"/> Passive motion</p> <p><input type="checkbox"/> On rest/non-movement    <input type="checkbox"/> Does not result in/cause functional loss</p> <p><input type="checkbox"/> Causes functional loss (if checked describe in the comments box below)</p>

**SECTION III - RANGE OF MOTION (ROM) AND FUNCTIONAL LIMITATION (continued)**

<p><b>3A. Initial ROM measurements (continued)</b></p> <p><b>Right shoulder</b></p> <p>Comments:</p> <div style="border: 1px solid black; height: 60px; margin-top: 5px;"></div> <p>Is there objective evidence of crepitus?   <input type="checkbox"/> Yes   <input checked="" type="checkbox"/> No</p> <p>Is there objective evidence of localized tenderness or pain on palpation of the joint or associated soft tissue?   <input type="checkbox"/> Yes   <input checked="" type="checkbox"/> No</p> <p>If yes, please explain. Include location, severity, and relationship to condition(s).</p> <div style="border: 1px solid black; height: 40px; margin-top: 5px;"></div>	<p><b>3A. Initial ROM measurements (continued)</b></p> <p><b>Left shoulder</b></p> <p>Comments:</p> <div style="border: 1px solid black; height: 60px; margin-top: 5px;"></div> <p>Is there objective evidence of crepitus?   <input type="checkbox"/> Yes   <input checked="" type="checkbox"/> No</p> <p>Is there objective evidence of localized tenderness or pain on palpation of the joint or associated soft tissue?   <input type="checkbox"/> Yes   <input checked="" type="checkbox"/> No</p> <p>If yes, please explain. Include location, severity, and relationship to condition(s).</p> <div style="border: 1px solid black; height: 40px; margin-top: 5px;"></div>
<p><b>3B. Observed repetitive use ROM</b></p> <p>Is the Veteran able to perform repetitive-use testing with at least three repetitions?</p> <p><input checked="" type="checkbox"/> Yes   <input type="checkbox"/> No</p> <p>If no, please explain:</p> <div style="border: 1px solid black; height: 60px; margin-top: 5px;"></div> <p>Is there additional loss of function or range of motion after three repetitions?</p> <p><input type="checkbox"/> Yes   <input checked="" type="checkbox"/> No</p> <p>If yes, please respond to the following after the completion of the three repetitions:</p> <p>Flexion endpoint (180 degrees): _____ degrees</p> <p>Abduction endpoint (180 degrees): _____ degrees</p> <p>Internal rotation endpoint (90 degrees): _____ degrees</p> <p>External rotation endpoint (90 degrees): _____ degrees</p> <p>Select factors that cause this functional loss (check all that apply):</p> <p><input type="checkbox"/> N/A   <input type="checkbox"/> Pain   <input type="checkbox"/> Fatigability   <input type="checkbox"/> Weakness</p> <p><input type="checkbox"/> Lack of endurance   <input type="checkbox"/> Incoordination</p> <p><input type="checkbox"/> Other _____</p>	<p><b>3B. Observed repetitive use ROM</b></p> <p>Is the Veteran able to perform repetitive-use testing with at least three repetitions?</p> <p><input checked="" type="checkbox"/> Yes   <input type="checkbox"/> No</p> <p>If no, please explain:</p> <div style="border: 1px solid black; height: 60px; margin-top: 5px;"></div> <p>Is there additional loss of function or range of motion after three repetitions?</p> <p><input type="checkbox"/> Yes   <input checked="" type="checkbox"/> No</p> <p>If yes, please respond to the following after the completion of the three repetitions:</p> <p>Flexion endpoint (180 degrees): _____ degrees</p> <p>Abduction endpoint (180 degrees): _____ degrees</p> <p>Internal rotation endpoint (90 degrees): _____ degrees</p> <p>External rotation endpoint (90 degrees): _____ degrees</p> <p>Select factors that cause this functional loss (check all that apply):</p> <p><input type="checkbox"/> N/A   <input type="checkbox"/> Pain   <input type="checkbox"/> Fatigability   <input type="checkbox"/> Weakness</p> <p><input type="checkbox"/> Lack of endurance   <input type="checkbox"/> Incoordination</p> <p><input type="checkbox"/> Other _____</p>
<p><small>Note: When pain is associated with movement, the examiner must give a statement on whether pain could significantly limit functional ability during flare-ups and/or after repeated use over time in terms of additional loss of range of motion. In the exam report, the examiner is requested to provide an estimate of decreased range of motion (in degrees) that reflect frequency, duration, and during flare-ups - even if not directly observed during a flare-up and/or after repeated use over time.</small></p>	
<p><b>3C. Repeated use over time</b></p> <p>Is the Veteran being examined immediately after repeated use over time?</p> <p><input checked="" type="checkbox"/> Yes   <input type="checkbox"/> No</p> <p>Does procured evidence (statements from the Veteran) suggest pain, fatigability, weakness, lack of endurance, or incoordination which significantly limits functional ability with repeated use over time?   <input type="checkbox"/> Yes   <input type="checkbox"/> No</p> <p>Select factors that cause this functional loss (check all that apply):</p> <p><input type="checkbox"/> N/A   <input type="checkbox"/> Pain   <input type="checkbox"/> Fatigability   <input type="checkbox"/> Weakness</p> <p><input type="checkbox"/> Lack of endurance   <input type="checkbox"/> Incoordination</p> <p><input type="checkbox"/> Other _____</p>	<p><b>3C. Repeated use over time</b></p> <p>Is the Veteran being examined immediately after repeated use over time?</p> <p><input checked="" type="checkbox"/> Yes   <input type="checkbox"/> No</p> <p>Does procured evidence (statements from the Veteran) suggest pain, fatigability, weakness, lack of endurance, or incoordination which significantly limits functional ability with repeated use over time?   <input type="checkbox"/> Yes   <input type="checkbox"/> No</p> <p>Select factors that cause this functional loss (check all that apply):</p> <p><input type="checkbox"/> N/A   <input type="checkbox"/> Pain   <input type="checkbox"/> Fatigability   <input type="checkbox"/> Weakness</p> <p><input type="checkbox"/> Lack of endurance   <input type="checkbox"/> Incoordination</p> <p><input type="checkbox"/> Other _____</p>



### SECTION III - RANGE OF MOTION (ROM) AND FUNCTIONAL LIMITATION (continued)

3C.Repeated use over time (continued)	3C.Repeated use over time (continued)
<p><b>Right shoulder</b></p> <p>Estimate range of motion in degrees for this joint immediately after repeated use over time based on information procured from relevant sources including the lay statements of the Veteran.</p> <p>Flexion endpoint (180 degrees): _____ degrees</p> <p>Abduction endpoint (180 degrees): _____ degrees</p> <p>Internal rotation endpoint (90 degrees): _____ degrees</p> <p>External rotation endpoint (90 degrees): _____ degrees</p> <p>The examiner should provide the estimated range of motion based on a review of all procurable information - to include the Veteran's statement on examination, case-specific evidence (to include medical treatment records when applicable and lay evidence), and the examiner's medical expertise. If, after evaluation of the procurable and assembled data, the examiner determines that it is not feasible to provide this estimate, the examiner should explain why an estimate cannot be provided. The explanation should not be based on an examiner's shortcomings or a general aversion to offering an estimate on issues not directly observed.</p> <p>Please cite and discuss evidence here. (Must be specific to the case and based on all procurable evidence.)</p>	<p><b>Left shoulder</b></p> <p>Estimate range of motion in degrees for this joint immediately after repeated use over time based on information procured from relevant sources including the lay statements of the Veteran.</p> <p>Flexion endpoint (180 degrees): _____ degrees</p> <p>Abduction endpoint (180 degrees): _____ degrees</p> <p>Internal rotation endpoint (90 degrees): _____ degrees</p> <p>External rotation endpoint (90 degrees): _____ degrees</p> <p>The examiner should provide the estimated range of motion based on a review of all procurable information - to include the Veteran's statement on examination, case-specific evidence (to include medical treatment records when applicable and lay evidence), and the examiner's medical expertise. If, after evaluation of the procurable and assembled data, the examiner determines that it is not feasible to provide this estimate, the examiner should explain why an estimate cannot be provided. The explanation should not be based on an examiner's shortcomings or a general aversion to offering an estimate on issues not directly observed.</p> <p>Please cite and discuss evidence here. (Must be specific to the case and based on all procurable evidence.)</p>
<p><b>3D. Flare-ups</b></p> <p>Is the examination being conducted during a flare-up?</p> <p><input type="checkbox"/> Yes    <input type="checkbox"/> No</p> <p>Does procured evidence (statements from the Veteran) suggest pain, fatigability, weakness, lack of endurance, or incoordination which significantly limits functional ability with flare-ups?    <input type="checkbox"/> Yes    <input type="checkbox"/> No</p> <p>Select factors that cause this functional loss (check all that apply):</p> <p><input type="checkbox"/> N/A    <input type="checkbox"/> Pain    <input type="checkbox"/> Fatigability    <input type="checkbox"/> Weakness</p> <p><input type="checkbox"/> Lack of endurance    <input type="checkbox"/> Incoordination</p> <p><input type="checkbox"/> Other _____</p> <p>Estimate range of motion in degrees for this joint during flare-ups based on information procured from relevant sources including the lay statements of the Veteran.</p> <p>Flexion endpoint (180 degrees): _____ degrees</p> <p>Abduction endpoint (180 degrees): _____ degrees</p> <p>Internal rotation endpoint (90 degrees): _____ degrees</p> <p>External rotation endpoint (90 degrees): _____ degrees</p> <p>The examiner should provide the estimated range of motion based on a review of all procurable information - to include the Veteran's statement on examination, case-specific evidence (to include medical treatment records when applicable and lay evidence), and the examiner's medical expertise. If, after evaluation of the procurable and assembled data, the examiner determines that it is not feasible to provide this estimate, the examiner should explain why an estimate cannot be provided. The explanation should not be based on an examiner's shortcomings or a general aversion to offering an estimate on issues not directly observed.</p> <p>Please cite and discuss evidence here. (Must be specific to the case and based on all procurable evidence.)</p>	<p><b>3D. Flare-ups</b></p> <p>Is the examination being conducted during a flare-up?</p> <p><input type="checkbox"/> Yes    <input type="checkbox"/> No</p> <p>Does procured evidence (statements from the Veteran) suggest pain, fatigability, weakness, lack of endurance, or incoordination which significantly limits functional ability with flare-ups?    <input type="checkbox"/> Yes    <input type="checkbox"/> No</p> <p>Select factors that cause this functional loss (check all that apply):</p> <p><input type="checkbox"/> N/A    <input type="checkbox"/> Pain    <input type="checkbox"/> Fatigability    <input type="checkbox"/> Weakness</p> <p><input type="checkbox"/> Lack of endurance    <input type="checkbox"/> Incoordination</p> <p><input type="checkbox"/> Other _____</p> <p>Estimate range of motion in degrees for this joint during flare-ups based on information procured from relevant sources including the lay statements of the Veteran.</p> <p>Flexion endpoint (180 degrees): _____ degrees</p> <p>Abduction endpoint (180 degrees): _____ degrees</p> <p>Internal rotation endpoint (90 degrees): _____ degrees</p> <p>External rotation endpoint (90 degrees): _____ degrees</p> <p>The examiner should provide the estimated range of motion based on a review of all procurable information - to include the Veteran's statement on examination, case-specific evidence (to include medical treatment records when applicable and lay evidence), and the examiner's medical expertise. If, after evaluation of the procurable and assembled data, the examiner determines that it is not feasible to provide this estimate, the examiner should explain why an estimate cannot be provided. The explanation should not be based on an examiner's shortcomings or a general aversion to offering an estimate on issues not directly observed.</p> <p>Please cite and discuss evidence here. (Must be specific to the case and based on all procurable evidence.)</p>

### SECTION III - RANGE OF MOTION (ROM) AND FUNCTIONAL LIMITATION (continued)

3E. Additional factors contributing to disability	3E. Additional factors contributing to disability
<p>In addition to those addressed above, are there additional contributing factors of disability? Select all that apply and describe:</p> <div style="display: flex; flex-wrap: wrap;"> <div style="width: 50%;"><input type="checkbox"/> None</div> <div style="width: 50%;"><input type="checkbox"/> Interference with sitting</div> <div style="width: 50%;"><input type="checkbox"/> Interference with standing</div> <div style="width: 50%;"><input type="checkbox"/> Swelling</div> <div style="width: 50%;"><input type="checkbox"/> Disturbance of locomotion</div> <div style="width: 50%;"><input type="checkbox"/> Deformity</div> <div style="width: 50%;"><input type="checkbox"/> Less movement than normal</div> <div style="width: 50%;"><input type="checkbox"/> More movement than normal</div> <div style="width: 50%;"><input type="checkbox"/> Weakened movement</div> <div style="width: 50%;"><input type="checkbox"/> Atrophy of disuse</div> <div style="width: 50%;"><input type="checkbox"/> Instability of station</div> <div style="width: 50%;"><input type="checkbox"/> Other, describe: <input style="width: 100%;" type="text"/></div> </div> <p>Please describe additional contributing factors of disability here:</p> <div style="border: 1px solid black; height: 100px; width: 100%;"></div>	<p>In addition to those addressed above, are there additional contributing factors of disability? Select all that apply and describe:</p> <div style="display: flex; flex-wrap: wrap;"> <div style="width: 50%;"><input type="checkbox"/> None</div> <div style="width: 50%;"><input type="checkbox"/> Interference with sitting</div> <div style="width: 50%;"><input type="checkbox"/> Interference with standing</div> <div style="width: 50%;"><input type="checkbox"/> Swelling</div> <div style="width: 50%;"><input type="checkbox"/> Disturbance of locomotion</div> <div style="width: 50%;"><input type="checkbox"/> Deformity</div> <div style="width: 50%;"><input type="checkbox"/> Less movement than normal</div> <div style="width: 50%;"><input type="checkbox"/> More movement than normal</div> <div style="width: 50%;"><input type="checkbox"/> Weakened movement</div> <div style="width: 50%;"><input type="checkbox"/> Atrophy of disuse</div> <div style="width: 50%;"><input type="checkbox"/> Instability of station</div> <div style="width: 50%;"><input type="checkbox"/> Other, describe: <input style="width: 100%;" type="text"/></div> </div> <p>Please describe additional contributing factors of disability here:</p> <div style="border: 1px solid black; height: 100px; width: 100%;"></div>

### SECTION IV - MUSCLE ATROPHY

Right shoulder	Left shoulder
<p>4A. Does the Veteran have muscle atrophy? <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>4B. If yes, is the muscle atrophy due to the claimed condition in the diagnosis section?  <input type="checkbox"/> Yes <input type="checkbox"/> No If no, provide rationale:</p> <div style="border: 1px solid black; height: 60px; width: 100%;"></div> <p>4C. For any muscle atrophy due to a diagnosis listed in Section I, indicate specific location of atrophy, providing measurements in centimeters of normal side and corresponding atrophied side, measured at maximum muscle bulk.</p> <p><input type="checkbox"/> Right upper extremity (specify location of measurement such as "10cm above the anterior elbow crease" here):</p> <div style="display: flex; justify-content: space-between; margin-top: 20px;"> <div>Circumference of more normal side: _____ cm</div> <div>Circumference of atrophied side: _____ cm</div> </div>	<p>4A. Does the Veteran have muscle atrophy? <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>4B. If yes, is the muscle atrophy due to the claimed condition in the diagnosis section?  <input type="checkbox"/> Yes <input type="checkbox"/> No If no, provide rationale:</p> <div style="border: 1px solid black; height: 60px; width: 100%;"></div> <p>4C. For any muscle atrophy due to a diagnosis listed in Section I, indicate specific location of atrophy, providing measurements in centimeters of normal side and corresponding atrophied side, measured at maximum muscle bulk.</p> <p><input type="checkbox"/> Left upper extremity (specify location of measurement such as "10cm above the anterior elbow crease" here):</p> <div style="display: flex; justify-content: space-between; margin-top: 20px;"> <div>Circumference of more normal side: _____ cm</div> <div>Circumference of atrophied side: _____ cm</div> </div>

### SECTION V - ANKYLOSIS

<p>Note: Ankylosis is the immobilization of a joint due to disease, injury, or surgical procedure.</p>	
<p>5A. Is there ankylosis of the scapulohumeral (glenohumeral) articulation (shoulder joint) - (i.e., the scapula and humerus move as one piece)? <input type="checkbox"/> Yes <input type="checkbox"/> No  If yes, indicate the severity of the ankylosis:</p> <div style="display: flex; flex-wrap: wrap;"> <div style="width: 50%;"><input type="checkbox"/> Ankylosis in abduction up to 60 degrees; can reach mouth and head (favorable ankylosis)</div> <div style="width: 50%;"><input type="checkbox"/> Ankylosis in abduction between favorable and unfavorable (intermediate ankylosis)</div> <div style="width: 50%;"><input type="checkbox"/> Ankylosis in abduction at 25 degrees or less from side (unfavorable ankylosis)</div> </div> <p>5B. Indicate angle of ankylosis in degrees of abduction: _____ degrees</p> <p>5C. If ankylosed, is there involvement of Muscle Group I (trapezius, levator scapulae, serratus magnus) and II (pectoralis major II (costosternal), latissimus dorsi and teres major, pectoralis minor; rhomboid)? <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, complete the Muscle Injuries questionnaire.</p>	<p>5A. Is there ankylosis of the scapulohumeral (glenohumeral) articulation (shoulder joint) - (i.e., the scapula and humerus move as one piece)? <input type="checkbox"/> Yes <input type="checkbox"/> No  If yes, indicate the severity of the ankylosis:</p> <div style="display: flex; flex-wrap: wrap;"> <div style="width: 50%;"><input type="checkbox"/> Ankylosis in abduction up to 60 degrees; can reach mouth and head (favorable ankylosis)</div> <div style="width: 50%;"><input type="checkbox"/> Ankylosis in abduction between favorable and unfavorable (intermediate ankylosis)</div> <div style="width: 50%;"><input type="checkbox"/> Ankylosis in abduction at 25 degrees or less from side (unfavorable ankylosis)</div> </div> <p>5B. Indicate angle of ankylosis in degrees of abduction: _____ degrees</p> <p>5C. If ankylosed, is there involvement of Muscle Group I (trapezius, levator scapulae, serratus magnus) and II (pectoralis major II (costosternal), latissimus dorsi and teres major, pectoralis minor; rhomboid)? <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, complete the Muscle Injuries questionnaire.</p>

## SECTION VI - ROTATOR CUFF CONDITIONS

<p>6A. Complete the following:</p> <p>Hawkins' Impingement Test: Forward flex the arm to 90 degrees with the elbow bent to 90 degrees. Internally rotate arm. Pain on internal rotation indicates a positive test; may signify rotator cuff tendinopathy or tear.</p> <p><input type="checkbox"/> Positive    <input type="checkbox"/> Negative    <input type="checkbox"/> Unable to test    <input type="checkbox"/> N/A</p> <p>Empty Can Test: Abduct arm to 90 degrees and forward flex 30 degrees. Patient turns thumbs down and resists downward force applied by the examiner. Weakness indicates a positive test; may indicate rotator cuff pathology, including supraspinatus tendinopathy or tear.</p> <p><input type="checkbox"/> Positive    <input type="checkbox"/> Negative    <input type="checkbox"/> Unable to test    <input type="checkbox"/> N/A</p> <p>External rotation/infraspinatus strength test: Patient holds arms at side with elbow flexed 90 degrees. Patient externally rotates against resistance. Weakness indicates a positive test; may be associated with infraspinatus tendinopathy or tear.</p> <p><input type="checkbox"/> Positive    <input type="checkbox"/> Negative    <input type="checkbox"/> Unable to test    <input type="checkbox"/> N/A</p> <p>Lift-off subscapularis test: Patient internally rotates arm behind lower back, pushes against examiner's hand. Weakness indicates a positive test; may indicate subscapularis tendinopathy or tear.</p> <p><input type="checkbox"/> Positive    <input type="checkbox"/> Negative    <input type="checkbox"/> Unable to test    <input type="checkbox"/> N/A</p> <p>6B. If unable to test, is a rotator cuff condition suspected? <input type="checkbox"/> Yes    <input type="checkbox"/> No If yes, please describe:</p> <div style="border: 1px solid black; height: 80px; margin-top: 5px;"></div>	<p>6A. Complete the following:</p> <p>Hawkins' Impingement Test: Forward flex the arm to 90 degrees with the elbow bent to 90 degrees. Internally rotate arm. Pain on internal rotation indicates a positive test; may signify rotator cuff tendinopathy or tear.</p> <p><input type="checkbox"/> Positive    <input type="checkbox"/> Negative    <input type="checkbox"/> Unable to test    <input type="checkbox"/> N/A</p> <p>Empty Can Test: Abduct arm to 90 degrees and forward flex 30 degrees. Patient turns thumbs down and resists downward force applied by the examiner. Weakness indicates a positive test; may indicate rotator cuff pathology, including supraspinatus tendinopathy or tear.</p> <p><input type="checkbox"/> Positive    <input type="checkbox"/> Negative    <input type="checkbox"/> Unable to test    <input type="checkbox"/> N/A</p> <p>External rotation/infraspinatus strength test: Patient holds arms at side with elbow flexed 90 degrees. Patient externally rotates against resistance. Weakness indicates a positive test; may be associated with infraspinatus tendinopathy or tear.</p> <p><input type="checkbox"/> Positive    <input type="checkbox"/> Negative    <input type="checkbox"/> Unable to test    <input type="checkbox"/> N/A</p> <p>Lift-off subscapularis test: Patient internally rotates arm behind lower back, pushes against examiner's hand. Weakness indicates a positive test; may indicate subscapularis tendinopathy or tear.</p> <p><input type="checkbox"/> Positive    <input type="checkbox"/> Negative    <input type="checkbox"/> Unable to test    <input type="checkbox"/> N/A</p> <p>6B. If unable to test, is a rotator cuff condition suspected? <input type="checkbox"/> Yes    <input type="checkbox"/> No If yes, please describe:</p> <div style="border: 1px solid black; height: 80px; margin-top: 5px;"></div>
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## SECTION VII - SHOULDER INSTABILITY, DISLOCATION OR LABRAL PATHOLOGY

<p><b>Right shoulder</b></p> <p>7A. Complete the following:</p> <p>Crank Apprehension and Relocation Test: With patient supine, abduct patient's arm to 90 degrees and flex elbow 90 degrees. Pain and sense of instability with further external rotation may indicate shoulder instability.</p> <p><input type="checkbox"/> Positive    <input type="checkbox"/> Negative    <input type="checkbox"/> Unable to test    <input type="checkbox"/> N/A</p> <p>7B. If unable to test, is shoulder instability, dislocation or labral pathology suspected? <input type="checkbox"/> Yes    <input type="checkbox"/> No If yes, please describe:</p> <div style="border: 1px solid black; height: 100px; margin-top: 5px;"></div> <p>7C. Is there shoulder instability, dislocation or labral pathology? <input type="checkbox"/> Yes    <input type="checkbox"/> No</p> <p>7D. Does the Veteran have mechanical symptoms (clicking, catching, etc.)? <input type="checkbox"/> Yes    <input type="checkbox"/> No</p> <p>7E. Are there current residuals of recurrent dislocation (subluxation) of the glenohumeral (scapulohumeral) joint? <input type="checkbox"/> Yes    <input type="checkbox"/> No    If yes, check all that apply:</p> <p><input type="checkbox"/> Infrequent episodes and guarding of movement only at shoulder level (flexion and/or abduction at 90°)</p> <p><input type="checkbox"/> Frequent episodes and guarding of all arm movements</p> <p>Affects range of motion? <input type="checkbox"/> Yes    <input type="checkbox"/> No</p>	<p><b>Left shoulder</b></p> <p>7A. Complete the following:</p> <p>Crank Apprehension and Relocation Test: With patient supine, abduct patient's arm to 90 degrees and flex elbow 90 degrees. Pain and sense of instability with further external rotation may indicate shoulder instability.</p> <p><input type="checkbox"/> Positive    <input type="checkbox"/> Negative    <input type="checkbox"/> Unable to test    <input type="checkbox"/> N/A</p> <p>7B. If unable to test, is shoulder instability, dislocation or labral pathology suspected? <input type="checkbox"/> Yes    <input type="checkbox"/> No If yes, please describe:</p> <div style="border: 1px solid black; height: 100px; margin-top: 5px;"></div> <p>7C. Is there shoulder instability, dislocation or labral pathology? <input type="checkbox"/> Yes    <input type="checkbox"/> No</p> <p>7D. Does the Veteran have mechanical symptoms (clicking, catching, etc.)? <input type="checkbox"/> Yes    <input type="checkbox"/> No</p> <p>7E. Are there current residuals of recurrent dislocation (subluxation) of the glenohumeral (scapulohumeral) joint? <input type="checkbox"/> Yes    <input type="checkbox"/> No    If yes, check all that apply:</p> <p><input type="checkbox"/> Infrequent episodes and guarding of movement only at shoulder level (flexion and/or abduction at 90°)</p> <p><input type="checkbox"/> Frequent episodes and guarding of all arm movements</p> <p>Affects range of motion? <input type="checkbox"/> Yes    <input type="checkbox"/> No</p>
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**SECTION VIII - CLAVICLE, SCAPULA, ACROMIOCLAVICULAR (AC) JOINT AND STERNOCLAVICULAR JOINT CONDITIONS**

8A. Complete the following:

Cross-body adduction test: Passively adduct arm across the patient's body toward the contralateral shoulder. Pain may indicate acromioclavicular joint pathology.

☐ Positive ☐ Negative ☐ Unable to test ☐ N/A8B. If unable to test, is a clavicle, scapula, acromioclavicular (AC) joint or sternoclavicular joint condition suspected? ☐ Yes ☐ No If yes, please describe:8C. Is there a clavicle, scapula, acromioclavicular (AC) joint, sternoclavicular joint condition or other impairment? ☐ Yes ☐ No If yes, indicate severity:

- ☐ Malunion of clavicle or scapula
- ☐ Nonunion of clavicle or scapula without loose movement
- ☐ Nonunion of clavicle or scapula with loose movement
- ☐ Dislocation (acromioclavicular separation or sternoclavicular dislocation)
- ☐ Other (describe):

8D. Does the clavicle or scapula condition affect range of motion of the shoulder (glenohumeral joint)? ☐ Yes ☐ No8E. Is there tenderness on palpation of the AC joint? ☐ Yes ☐ No

8A. Complete the following:

Cross-body adduction test: Passively adduct arm across the patient's body toward the contralateral shoulder. Pain may indicate acromioclavicular joint pathology.

☐ Positive ☐ Negative ☐ Unable to test ☐ N/A8B. If unable to test, is a clavicle, scapula, acromioclavicular (AC) joint or sternoclavicular joint condition suspected? ☐ Yes ☐ No If yes, please describe:8C. Is there a clavicle, scapula, acromioclavicular (AC) joint, sternoclavicular joint condition or other impairment? ☐ Yes ☐ No If yes, indicate severity:

- ☐ Malunion of clavicle or scapula
- ☐ Nonunion of clavicle or scapula without loose movement
- ☐ Nonunion of clavicle or scapula with loose movement
- ☐ Dislocation (acromioclavicular separation or sternoclavicular dislocation)
- ☐ Other (describe):

8D. Does the clavicle or scapula condition affect range of motion of the shoulder (glenohumeral joint)? ☐ Yes ☐ No8E. Is there tenderness on palpation of the AC joint? ☐ Yes ☐ No**SECTION IX - CONDITIONS OR IMPAIRMENTS OF THE HUMERUS**9A. Does the Veteran have loss of head (flail shoulder), nonunion (false flail shoulder), or fibrous union of the humerus? ☐ Yes ☐ No If yes, check all that apply:☐ Loss of head (flail shoulder) ☐ Nonunion (false flail shoulder) ☐ Fibrous union9B. Does the Veteran have malunion of the humerus with moderate or marked deformity?: ☐ Yes ☐ No If yes, indicate severity:☐ Moderate deformity ☐ Marked deformity9C. Does the humerus condition affect range of motion of the shoulder (glenohumeral joint)? ☐ Yes ☐ No9A. Does the Veteran have loss of head (flail shoulder), nonunion (false flail shoulder), or fibrous union of the humerus? ☐ Yes ☐ No If yes, check all that apply:☐ Loss of head (flail shoulder) ☐ Nonunion (false flail shoulder) ☐ Fibrous union9B. Does the Veteran have malunion of the humerus with moderate or marked deformity?: ☐ Yes ☐ No If yes, indicate severity:☐ Moderate deformity ☐ Marked deformity9C. Does the humerus condition affect range of motion of the shoulder (glenohumeral joint)? ☐ Yes ☐ No**SECTION X - SURGICAL PROCEDURES**

10. Indicate any surgical procedures that the Veteran has had performed and provide the additional information as requested (check all that apply):

☐ No surgery☐ Total shoulder joint replacement Date of surgery: \_\_\_\_\_Residuals: ☐ None ☐ Intermediate degrees of residual weakness, pain, or limitation of motion☐ Chronic residuals consisting of severe painful motion or weakness☐ Other residuals, describe: \_\_\_\_\_☐ Arthroscopic or other shoulder surgery

Date of Surgery: \_\_\_\_\_ Type of Surgery: \_\_\_\_\_

Describe residuals:

10. Indicate any surgical procedures that the Veteran has had performed and provide the additional information as requested (check all that apply):

☐ No surgery☐ Total shoulder joint replacement Date of surgery: \_\_\_\_\_Residuals: ☐ None ☐ Intermediate degrees of residual weakness, pain, or limitation of motion☐ Chronic residuals consisting of severe painful motion or weakness☐ Other residuals, describe: \_\_\_\_\_☐ Arthroscopic or other shoulder surgery

Date of Surgery: \_\_\_\_\_ Type of Surgery: \_\_\_\_\_

Describe residuals:

**SECTION XI - OTHER PERTINENT PHYSICAL FINDINGS, COMPLICATIONS, CONDITIONS, SIGNS, SYMPTOMS, AND SCARS**

11A. Does the Veteran have any other pertinent physical findings, complications, signs, or symptoms related to any conditions listed in the diagnosis section above?

☐ Yes ☐ No If yes, describe (brief summary):

11B. 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 the appropriate dermatological questionnaire.

11C. Comments, if any:

**SECTION XII - ASSISTIVE DEVICES**

12A. Does the Veteran use any assistive devices? ☐ Yes ☐ No

If yes, identify the assistive devices used. Check all that apply and indicate frequency:

☐ Brace Frequency of use: ☐ Occasional ☐ Regular ☐ Constant

☐ Other, describe: \_\_\_\_\_ Frequency of use: ☐ Occasional ☐ Regular ☐ Constant

12B. If the Veteran uses any assistive devices, specify the condition, indicate the side, and identify the assistive device used for each condition:

**SECTION XIII - REMAINING EFFECTIVE FUNCTION OF THE EXTREMITIES**

Note: The intention of this section is to permit the examiner to quantify the level of remaining function; it is not intended to inquire whether the Veteran should undergo an amputation with fitting of a prosthesis. For example, if the functions of grasping (hand) or propulsion (foot) are as limited as if the Veteran had an amputation and prosthesis, the examiner should check "yes" and describe the diminished functioning. The question simply asks whether the functional loss is to the same degree as if there were an amputation of the affected limb.

13A. Due to the Veteran's shoulder or arm condition(s), is there functional impairment of an extremity such that no effective functions remain other than that which would be equally well-served by an amputation with prosthesis (functions of the upper extremity include grasping, manipulation, etc.)?

☐ Yes, functioning is so diminished that amputation with prosthesis would equally serve the Veteran

☐ No

If yes, indicate extremities for which this applies: ☐ Right upper ☐ Left upper

13B. For each checked extremity, identify the condition causing loss of function, describe loss of effective function, and provide specific examples (brief summary):

**SECTION XIV - DIAGNOSTIC TESTING**

Note: Testing listed below is not indicated for every condition. The diagnosis of degenerative arthritis (osteoarthritis) or post-traumatic arthritis must be confirmed by imaging studies. Once such arthritis has been documented, even if in the past, no further imaging studies are required by VA, even if arthritis has worsened.

14A. Have imaging studies been performed in conjunction with this examination? ☒ Yes ☐ No

14B. If yes, is degenerative or post-traumatic arthritis documented? ☐ Yes ☒ No If yes, indicate side: ☐ Right ☐ Left ☐ Both

14C. If yes, provide type of test or procedure, date and results (brief summary):

**SECTION XIV - DIAGNOSTIC TESTING (continued)**

14D. Are there any other significant diagnostic test findings or results related to the claimed condition(s) and/or diagnosis(es), that were reviewed in conjunction with this examination?  
☐ Yes ☐ No If yes, provide type of test or procedure, date and results (brief summary):

14E. If any test results are other than normal, indicate relationship of abnormal findings to diagnosed condition(s):

**SECTION XV - FUNCTIONAL IMPACT**

Note: Provide the impact of only the diagnosed condition(s), without consideration of the impact of other medical conditions or factors, such as age.

15A. Regardless of the Veteran's current employment status, do the conditions listed in the diagnosis section impact his/her ability to perform any type of occupational task (such as standing, walking, lifting, sitting, etc.)? ☐ Yes ☐ No If yes, describe the functional impact of each condition, providing one or more examples:

**SECTION XVI - REMARKS**

16A. Remarks (if any – please identify the section to which the remark pertains when appropriate):

For issue of right shoulder pain refer to diagnosis section

**SECTION XVII - EXAMINER'S CERTIFICATION AND SIGNATURE**

**CERTIFICATION** - To the best of my knowledge, the information contained herein is accurate, complete and current.

17A. Examiner's signature

17B. Examiner's printed name:

17C. Date signed

Robert Smith NP

08/26/2020

17D. Examiner's phone/fax numbers

17E. National Provider Identifier (NPI) number

17F. Medical license number and state

619-400-1234

123456

123456

17G. Examiner's address

8810 rio san diego dr, san diego, ca 92108

**INTERNAL VETERANS AFFAIRS USE  
HEARING LOSS AND TINNITUS  
DISABILITY BENEFITS QUESTIONNAIRE**

**IMPORTANT -** THE DEPARTMENT OF VETERANS AFFAIRS (VA) ***WILL NOT PAY OR REIMBURSE*** ANY EXPENSES OR COST INCURRED IN THE PROCESS OF COMPLETING AND/OR SUBMITTING THIS FORM. PLEASE READ THE PRIVACY ACT AND RESPONDENT BURDEN INFORMATION BEFORE COMPLETING FORM.

NAME OF PATIENT/VETERAN

John James Rambo

PATIENT/VETERAN'S SOCIAL SECURITY NUMBER

xxx-xx-0012

Your patient is applying to the U. S. Department of Veterans Affairs (VA) for disability benefits. VA will consider the information you provide on this questionnaire as part of their evaluation in processing the Veteran's claim. Please note that this questionnaire is for disability evaluation, not for treatment purposes.

IS THIS QUESTIONNAIRE BEING COMPLETED IN CONJUNCTION WITH A VA21-2507, C&P EXAMINATION REQUEST?

☒ YES ☐ NO

How was the examination completed? (check all that apply)

- ☒ In-person examination  
☐ Records reviewed  
☐ Examination via approved video telehealth  
☐ Other, please specify in comments box:

Comments:

**ACCEPTABLE CLINICAL EVIDENCE (ACE)**

INDICATE METHOD USED TO OBTAIN MEDICAL INFORMATION TO COMPLETE THIS DOCUMENT:

- ☐ Review of available records (without in-person or video telehealth examination) using the Acceptable Clinical Evidence (ACE) process because the existing medical evidence provided sufficient information on which to prepare the questionnaire and such an examination will likely provide no additional relevant evidence.
- ☐ Review of available records in conjunction with an interview with the Veteran (without in-person or telehealth examination) using the ACE process because the existing medical evidence supplemented with an interview provided sufficient information on which to prepare the questionnaire and such an examination would likely provide no additional relevant evidence.

**EVIDENCE REVIEW**

EVIDENCE REVIEWED (check all that apply):

- ☐ Not requested ☐ No records were reviewed  
☐ VA claims file (hard copy paper C-file)  
☐ VA e-folder (VBMS or Virtual VA)  
☐ CPRS  
☐ Other (please identify other evidence reviewed):

EVIDENCE COMMENTS:

**NOTE:** This form is only for use by VHA staff or contract examiners.

This exam is for:

- ☐ Tinnitus only (audiologist or non-audiologist clinician) **If this exam is for tinnitus only, complete section 2 only. Otherwise complete entire form.**  
☒ Hearing loss and/or tinnitus (audiologist, performing current exam)  
☐ Hearing loss and/or tinnitus (audiologist or non-audiologist clinician, using audiology report of record that represents Veteran's current condition)

If using audiology report of record, date audiology exam was performed:

**SECTION 1: HEARING LOSS (HL)**

**Note: All testing must be conducted in accordance with the following instructions to be valid for VA disability evaluation purposes.**

**Instructions:** An examination of hearing impairment must be conducted by a state-licensed audiologist and must include a controlled speech discrimination test (specifically, the Maryland CNC recording) and a puretone audiometry test in a sound isolated booth that meets American National Standards Institute standards (ANSI S3.1.1999 [R2004]) for ambient noise. Measurements will be reported at the frequencies of 500, 1000, 2000, 3000, and 4000 Hz.

The examination will include the following tests: Puretone audiometry by air conduction at 250, 500, 1000, 2000, 3000, 4000, 6000 Hz and 8000 Hz, and by bone conduction at 250, 500, 1000, 2000, 3000, and 4000 Hz, spondee thresholds, speech discrimination using the recorded Maryland CNC Test, tympanometry and acoustic reflex tests (ipsilateral and contralateral), and, when necessary, Stenger tests. Bone conduction thresholds are measured when the air conduction thresholds are poorer than 15 dB HL. A modified Hughson-Westlake procedure will be used with appropriate masking. A Stenger must be administered whenever puretone air conduction thresholds at 500, 1000, 2000, 3000, and 4000 Hz differ by 20 dB or more between the two ears.

Maximum speech discrimination will be reported with the 50 word VA approved recording of the Maryland CNC test. The starting presentation level will be 40 dB re SRT. If necessary, the starting level will be adjusted upward to obtain a level at least 5 dB above the threshold at 2000 Hz, if not above the patient's tolerance level.

The examination will be conducted without the use of hearing aids. Both ears must be examined for hearing impairment even if hearing loss in only one ear is at issue.

When speech discrimination is 92% or less, a performance intensity function must be obtained.

A comprehensive audiological evaluation should include evaluation results for puretone thresholds by air and bone conduction (500-8000 Hz), speech reception thresholds (SRT), speech discrimination scores, and acoustic immittance with acoustic reflexes (ipsilateral and contralateral reflexes). Tests for non-organicity must be performed when indicated.

**1. OBJECTIVE FINDINGS****A. PURETONE THRESHOLDS IN DECIBELS (AIR CONDUCTION):**

Instructions: Measure and record puretone threshold values in decibels at the indicated frequencies (air conduction). Report the decibel (dB) value, which ranges from -10 dB to 105 dB, for each of the frequencies. Add a plus behind the decibel value when a maximum value has been reached with a failure of response from the Veteran. In those circumstances where the average includes a failure of response at either the maximum allowable limit (105 dB) or the maximum limits of the audiometer, use this maximum decibel value of the failure of response in the puretone threshold average calculation.

If the Veteran could not be tested (CNT), enter CNT and state the reason why the Veteran could not be tested. Clearly inaccurate, invalid or unreliable test results should not be reported.

The puretone threshold at 500 Hz is not used in calculating the puretone threshold average for evaluation purposes but is used in determining whether or not for VA purposes, hearing impairment reaches the level of a disability. The puretone threshold average requires the decibel levels of each of the required frequencies (1000 Hz, 2000 Hz, 3000 Hz, and 4000 Hz) be recorded for the test to be valid for determination of a hearing impairment.

**RIGHT EAR**

A	B	C	D	E	F	G	
500 Hz*	1000 Hz*	2000 Hz*	3000 Hz*	4000 Hz*	6000 Hz*	8000 Hz*	Avg Hz (B-E)**
15	15	25	25	25	20	25	22.5

**LEFT EAR**

A	B	C	D	E	F	G	
500 Hz*	1000 Hz*	2000 Hz*	3000 Hz*	4000 Hz*	6000 Hz*	8000 Hz*	Avg Hz (B-E)**
20	15	25	25	30	20	35	23.75

\*The puretone threshold at 500 Hz is not used in determining the evaluation but is used in determining whether or not a ratable hearing loss exists.

\*\*The average of B, C, D, and E.

\*\*\*CNT - Could Not Test

**B. WERE THERE ONE OR MORE FREQUENCY(IES) THAT COULD NOT BE TESTED?**

☐ YES ☒ NO *If yes, enter CNT in the box for frequency(ies) that could not be tested, and explain why testing could not be done:*

**C. VALIDITY OF PURETONE TEST RESULTS:**

- ☒ Test results are valid for rating purposes.
- ☐ Test results are not valid for rating purposes (not indicative of organic hearing loss).
- If invalid, provide reason:

**D. SPEECH DISCRIMINATION SCORE (MARYLAND CNC WORD LIST)**

Instructions on pausing: Examiners should pause when necessary during speech discrimination tests, in order to give the Veteran sufficient time to respond. This will ensure that the test results are based on actual hearing loss rather than on the effects of other problems that might slow a Veteran's response. There are a variety of problems that might require pausing, for example, the presence of cognitive impairment. It is up to the examiner to determine when to use pausing and the length of the pauses.

RIGHT EAR	100	%
LEFT EAR	100	%



**E. APPROPRIATENESS OF USE OF WORD RECOGNITION SCORE (MARYLAND CNC WORD LIST):**

RIGHT EAR:

IS WORD DISCRIMINATION SCORE AVAILABLE?

☒ YES ☐ NO☒ Use of speech discrimination score is appropriate for this Veteran.☐ The use of the speech discrimination score is not appropriate for this Veteran because of language difficulties, cognitive problems, inconsistent speech discrimination scores, etc., that make combined use of puretone average and speech discrimination scores inappropriate.

LEFT EAR:

IS WORD DISCRIMINATION SCORE AVAILABLE?

☒ YES ☐ NO☒ Use of speech discrimination score is appropriate for this Veteran.☐ The use of the speech discrimination score is not appropriate for this Veteran because of language difficulties, cognitive problems, inconsistent speech discrimination scores, etc., that make combined use of puretone average and speech discrimination scores inappropriate.**F. AUDIOLOGIC FINDINGS**

Summary of Immittance (Tympanometry) Findings:

	RIGHT EAR		LEFT EAR	
ACOUSTIC IMMITTANCE	<input checked="" type="checkbox"/> Normal	<input type="checkbox"/> Abnormal	<input checked="" type="checkbox"/> Normal	<input type="checkbox"/> Abnormal
IPSILATERAL ACOUSTIC REFLEXES	<input checked="" type="checkbox"/> Normal	<input type="checkbox"/> Abnormal	<input checked="" type="checkbox"/> Normal	<input type="checkbox"/> Abnormal
CONTRALATERAL ACOUSTIC REFLEXES	<input checked="" type="checkbox"/> Normal	<input type="checkbox"/> Abnormal	<input checked="" type="checkbox"/> Normal	<input type="checkbox"/> Abnormal
UNABLE TO INTERPRET REFLEXES DUE TO ARTIFACT	<input type="checkbox"/>		<input type="checkbox"/>	
UNABLE TO OBTAIN / MAINTAIN SEAL	<input type="checkbox"/>		<input type="checkbox"/>	

**2. DIAGNOSIS**RIGHT EAR

<input checked="" type="checkbox"/> Normal hearing	ICD CODE:
<input type="checkbox"/> Conductive hearing loss	ICD CODE:
<input type="checkbox"/> Mixed hearing loss	ICD CODE:
<input type="checkbox"/> Sensorineural hearing loss (in frequency range of 500-4000 Hz)*	ICD CODE:
<input type="checkbox"/> Sensorineural hearing loss (in frequency range of 6000 Hz or higher frequencies)**	ICD CODE:
<input type="checkbox"/> Significant changes in hearing thresholds in service***	ICD CODE:

LEFT EAR

<input type="checkbox"/> Normal hearing	ICD CODE:
<input type="checkbox"/> Conductive hearing loss	ICD CODE:
<input type="checkbox"/> Mixed hearing loss	ICD CODE:
<input checked="" type="checkbox"/> Sensorineural hearing loss (in frequency range of 500-4000 Hz)*	ICD CODE: H90
<input checked="" type="checkbox"/> Sensorineural hearing loss (in frequency range of 6000 Hz or higher frequencies)**	ICD CODE: H90
<input type="checkbox"/> Significant changes in hearing thresholds in service***	

**NOTES:**

\*The Veteran may have hearing loss at a level that is not considered to be a disability for VA purposes. This can occur when the auditory thresholds are greater than 25 dB at one or more frequencies in the 500-4000 Hz range.

\*\* The Veteran may have impaired hearing, but it does not meet the criteria to be considered a disability for VA purposes. For VA purposes, the diagnosis of hearing impairment is based upon testing at frequency ranges of 500, 1000, 2000, 3000, and 4000 Hz. If there is no HL in the 500-4000 Hz range, but there is HL above 4000 Hz, check this box.

\*\*\*The Veteran may have a significant change in hearing threshold in service, but it does not meet the criteria to be considered a disability for VA purposes. (A significant change in hearing threshold may indicate noise exposure or acoustic trauma.)

**3. ETIOLOGY**☐ ETIOLOGY OPINION NOT INDICATED AS: ☐ SERVICE CONNECTED CONDITION ☐ VBA DID NOT REQUEST ETIOLOGYRIGHT EAR

WAS THERE A PERMANENT POSITIVE THRESHOLD SHIFT (WORSE THAN REFERENCE THRESHOLD) GREATER THAN NORMAL MEASUREMENT VARIABILITY AT ANY FREQUENCY BETWEEN 500 AND 6000 HZ FOR THE RIGHT EAR?

☐ YES ☒ NO

OPINION PROVIDED FOR THE RIGHT EAR:

☐ YES ☒ NO

### 3. ETIOLOGY (continued)

#### RIGHT EAR (continued)

IF PRESENT, IS THE VETERAN'S RIGHT EAR HEARING LOSS AT LEAST AS NOT (50% PROBABILITY OR GREATER) CAUSED BY OR A RESULT OF AN EVENT IN MILITARY SERVICE?

☐ YES

☐ NO

☐ CANNOT DETERMINE A MEDICAL OPINION REGARDING THE ETIOLOGY OF THE VETERAN'S RIGHT EAR HEARING LOSS WITHOUT RESORTING TO SPECULATION:

RATIONALE (Provide rationale for either a yes, no answer or speculation reason):

DID HEARING LOSS EXIST PRIOR TO SERVICE?

☐ YES ☐ NO

IF YES, WAS THE PRE-EXISTING HEARING LOSS AGGRAVATED BEYOND NORMAL PROGRESSION IN MILITARY SERVICE?

☐ YES ☐ NO

PROVIDE RATIONALE FOR BOTH YES OR NO:

#### LEFT EAR

WAS THERE A PERMANENT POSITIVE THRESHOLD SHIFT (WORSE THAN REFERENCE THRESHOLD) GREATER THAN NORMAL MEASUREMENT VARIABILITY AT ANY FREQUENCY BETWEEN 500 AND 6000 HZ FOR THE LEFT EAR?

☐ YES ☒ NO

OPINION PROVIDED FOR THE LEFT EAR:

☐ YES ☒ NO

IF PRESENT, IS THE VETERAN'S LEFT EAR HEARING LOSS AT LEAST AS NOT (50% PROBABILITY OR GREATER) CAUSED BY OR A RESULT OF AN EVENT IN MILITARY SERVICE?

☐ YES

☐ NO

☐ CANNOT DETERMINE A MEDICAL OPINION REGARDING THE ETIOLOGY OF THE VETERAN'S LEFT EAR HEARING LOSS WITHOUT RESORTING TO SPECULATION:

RATIONALE (Provide rationale for either a yes, no answer or speculation reason):

DID HEARING LOSS EXIST PRIOR TO SERVICE?

☐ YES ☒ NO

IF YES, WAS THE PRE-EXISTING HEARING LOSS AGGRAVATED BEYOND NORMAL PROGRESSION IN MILITARY SERVICE?

☐ YES ☐ NO

PROVIDE RATIONALE FOR BOTH YES OR NO:

### 4. FUNCTIONAL IMPACT OF HEARING LOSS

NOTE: Ask the Veteran to describe in his or her own words the effects of disability (i.e., the current complaint of hearing loss on occupational functioning and daily activities). Document the Veteran's response without opining on the relationship between the functional effects and the level of impairment (audiogram) or otherwise characterizing the response. Do not use handicap scales.

DOES THE VETERAN'S HEARING LOSS IMPACT ORDINARY CONDITIONS OF DAILY LIFE, INCLUDING ABILITY TO WORK?

☒ YES ☐ NO

IF YES, DESCRIBE IMPACT IN THE VETERAN'S OWN WORDS:

Difficulty understanding those around me. Required to stand close to understand.  
Still on active duty as special forces.

**5. REMARKS, IF ANY, PERTAINING TO HEARING LOSS:**

Pre-military: None During-military: M9 handgun, M16 rifle, M4 Rifle, M249 Rifle, hand grenade training, right handed shooter. 30+ years as a active duty special forces. aircraft.

The examinee reports serving in the Army. The examinee indicates they served a total of 40+ year(s). The period(s) of service were from: 11/17/1965 to 7/31/2020. The examinee was in service during Vietnam War, Gulf War, Afghan War and Iraq War. The examinee reports that they participated in combat activity.

For the claimant's claimed condition of hearing loss, left please refer to the diagnosis

**SECTION 2: TINNITUS****1. MEDICAL HISTORY**

DOES THE VETERAN REPORT RECURRENT TINNITUS?

☒ YES ☐ NO

DATE AND CIRCUMSTANCES OF ONSET OF TINNITUS:

Approximately 20 years ago after a night of flying on a C130 I noticed a ringing in my ears while trying to go to sleep. The reported tinnitus is constant. The side(s) affected:

Both

**2. ETIOLOGY OF TINNITUS**

SELECT ANSWER BELOW AND PROVIDE RATIONALE WHERE REQUESTED:

- ☐ ETIOLOGY OPINION NOT INDICATED AS: ☐ SERVICE CONNECTED CONDITION ☐ VBA DID NOT REQUEST ETIOLOGY
- ☐ THE VETERAN HAS A DIAGNOSIS OF CLINICAL HEARING LOSS, AND HIS OR HER TINNITUS IS AT LEAST AS LIKELY AS NOT (50% PROBABILITY OR GREATER) A SYMPTOM ASSOCIATED WITH THE HEARING LOSS, AS TINNITUS IS KNOWN TO BE A SYMPTOM ASSOCIATED WITH HEARING LOSS.
- ☐ LESS LIKELY THAN NOT (LESS THAN 50% PROBABILITY) A SYMPTOM ASSOCIATED WITH THE VETERAN'S HEARING LOSS

RATIONALE:

- ☐ AT LEAST AS LIKELY AS NOT (50% PROBABILITY OR GREATER) CAUSED BY OR A RESULT OF MILITARY NOISE EXPOSURE

RATIONALE:

- ☐ AT LEAST AS LIKELY AS NOT (50% PROBABILITY OR GREATER) DUE TO A KNOWN ETIOLOGY (*such as traumatic brain injury*)

RATIONALE:

- ☐ LESS LIKELY THAN NOT (LESS THAN 50% PROBABILITY) CAUSED BY OR A RESULT OF MILITARY NOISE EXPOSURE

RATIONALE:

- ☒ CANNOT PROVIDE A MEDICAL OPINION REGARDING THE ETIOLOGY OF THE VETERAN'S TINNITUS WITHOUT RESORTING TO SPECULATION

REASON SPECULATION REQUIRED:

See remarks

### 3. FUNCTIONAL IMPACT OF TINNITUS

NOTE: Ask the Veteran to describe in his or her own words the effects of disability (i.e., the current complaint on occupational functioning and daily activities). Document the Veteran's response without opining on the relationship between the functional effects and the level of impairment (audiogram) or otherwise characterizing the response. Do not use handicap scales.

DOES THE VETERAN'S TINNITUS IMPACT ORDINARY CONDITIONS OF DAILY LIFE, INCLUDING ABILITY TO WORK?

☒ YES ☐ NO

IF YES, DESCRIBE IMPACT IN THE VETERAN'S OWN WORDS

Ringling makes it difficult to focus for sustained periods.

### 4. REMARKS, IF ANY, PERTAINING TO TINNITUS

For issue of hearing loss please refer to diagnosis section.

For the issue of Tinnitus, I am unable to provide diagnosis without mere speculation.

### SECTION 3: PHYSICIAN'S CERTIFICATION AND SIGNATURE

**CERTIFICATION** - To the best of my knowledge, the information contained herein is accurate, complete and current.

3A. AUDIOLOGIST/PHYSICIAN SIGNATURE & TITLE	3B. AUDIOLOGIST/PHYSICIAN PRINTED NAME  Richard Jones Nurse Practitioner
3C. DATE SIGNED	3D. AUDIOLOGIST/PHYSICIAN PHONE AND FAX NUMBER 619-400-1234
3E. NATIONAL PROVIDER IDENTIFIER (NPI) NUMBER  NPI: 12566666 Lic#:A#697 AR	3F. AUDIOLOGIST/PHYSICIAN ADDRESS  8810 Rio San Diego Drive, San Diego, CA 92108

**NOTE** - VA may request additional medical information, including additional examinations, if necessary to complete VA's review of the veteran's application.

**IMPORTANT** - Audiologist/Physician please fax the completed form to \_\_\_\_\_

(VA Regional Office FAX No.)

**NOTE** - A list of VA Regional Office FAX Numbers can be found at [www.benefits.va.gov/disabilityexams](http://www.benefits.va.gov/disabilityexams) or obtained by calling 1-800-827-1000.

**PRIVACY ACT NOTICE:** VA will not disclose information collected on this form to any source other than what has been authorized under the Privacy Act of 1974 or Title 38, Code of Federal Regulations 1.576 for routine uses (i.e., civil or criminal law enforcement, congressional communications, epidemiological or research studies, the collection of money owed to the United States, litigation in which the United States is a party or has an interest, the administration of VA programs and delivery of VA benefits, verification of identity and status, and personnel administration) as identified in the VA system of records, 58/VA21/22/28, Compensation, Pension, Education and Vocational Rehabilitation and Employment Records - VA, published in the Federal Register. Your obligation to respond is voluntary. VA uses your SSN to identify your claim file. Providing your SSN will help ensure that your records are properly associated with your claim file. Giving us your SSN account information is voluntary. Refusal to provide your SSN by itself will not result in the denial of benefits. VA will not deny an individual benefits for refusing to provide his or her SSN unless the disclosure of the SSN is required by a Federal Statute of law in effect prior to January 1, 1975, and still in effect. The requested information is considered relevant and necessary to determine maximum benefits under the law. The responses you submit are considered confidential (38 U.S.C. 5701). Information submitted is subject to verification through computer matching programs with other agencies.

**RESPONDENT BURDEN:** We need this information to determine entitlement to benefits (38 U.S.C. 501). Title 38, United States Code, allows us to ask for this information. We estimate that you will need an average of 30 minutes to review the instructions, find the information, and complete the form. VA cannot conduct or sponsor a collection of information unless a valid OMB control number is displayed. You are not required to respond to a collection of information if this number is not displayed. Valid OMB control numbers can be located on the OMB Internet Page at [www.reginfo.gov/public/do/PRAMain](http://www.reginfo.gov/public/do/PRAMain). If desired, you can call 1-800-827-1000 to get information on where to send comments or suggestions about this form.

**INTERNAL VETERANS AFFAIRS USE  
HEARING LOSS AND TINNITUS  
DISABILITY BENEFITS QUESTIONNAIRE**

**IMPORTANT -** THE DEPARTMENT OF VETERANS AFFAIRS (VA) ***WILL NOT PAY OR REIMBURSE*** ANY EXPENSES OR COST INCURRED IN THE PROCESS OF COMPLETING AND/OR SUBMITTING THIS FORM. PLEASE READ THE PRIVACY ACT AND RESPONDENT BURDEN INFORMATION BEFORE COMPLETING FORM.

NAME OF PATIENT/VETERAN John James Rambo	PATIENT/VETERAN'S SOCIAL SECURITY NUMBER xxx-xx-0012
---------------------------------------------	---------------------------------------------------------

Your patient is applying to the U. S. Department of Veterans Affairs (VA) for disability benefits. VA will consider the information you provide on this questionnaire as part of their evaluation in processing the Veteran's claim. Please note that this questionnaire is for disability evaluation, not for treatment purposes.

IS THIS QUESTIONNAIRE BEING COMPLETED IN CONJUNCTION WITH A VA21-2507, C&P EXAMINATION REQUEST?

☒ YES    ☐ NO

How was the examination completed? (check all that apply)

- ☒ In-person examination  
☒ Records reviewed  
☒ Examination via approved video telehealth  
☐ Other, please specify in comments box:

Comments:

**ACCEPTABLE CLINICAL EVIDENCE (ACE)**

INDICATE METHOD USED TO OBTAIN MEDICAL INFORMATION TO COMPLETE THIS DOCUMENT:

- ☐ Review of available records (without in-person or video telehealth examination) using the Acceptable Clinical Evidence (ACE) process because the existing medical evidence provided sufficient information on which to prepare the questionnaire and such an examination will likely provide no additional relevant evidence.
- ☐ Review of available records in conjunction with an interview with the Veteran (without in-person or telehealth examination) using the ACE process because the existing medical evidence supplemented with an interview provided sufficient information on which to prepare the questionnaire and such an examination would likely provide no additional relevant evidence.

**EVIDENCE REVIEW**

EVIDENCE REVIEWED (check all that apply):

- |                                                                           |                                                   |
|---------------------------------------------------------------------------|---------------------------------------------------|
| <input type="checkbox"/> Not requested                                    | <input type="checkbox"/> No records were reviewed |
| <input type="checkbox"/> VA claims file (hard copy paper C-file)          |                                                   |
| <input checked="" type="checkbox"/> VA e-folder (VBMS or Virtual VA)      |                                                   |
| <input type="checkbox"/> CPRS                                             |                                                   |
| <input type="checkbox"/> Other (please identify other evidence reviewed): |                                                   |

EVIDENCE COMMENTS:

All pertinent information was reviewed. Audiogram dated 2000 and last audiogram on file dated 2018 show hearing within normal limits bilaterally. Veteran is currently in the Army serving with an MOS of special forces, which has a high probability of hazardous noise exposure. In addition, he fought in combat during the Vietnam War.

**NOTE:** This form is only for use by VHA staff or contract examiners.

This exam is for:

- ☐ Tinnitus only (audiologist or non-audiologist clinician)    If this exam is for tinnitus only, complete section 2 only. Otherwise complete entire form.
- ☒ Hearing loss and/or tinnitus (audiologist, performing current exam)
- ☐ Hearing loss and/or tinnitus (audiologist or non-audiologist clinician, using audiology report of record that represents Veteran's current condition)

If using audiology report of record, date audiology exam was performed:

**SECTION 1: HEARING LOSS (HL)**

**Note: All testing must be conducted in accordance with the following instructions to be valid for VA disability evaluation purposes.**

**Instructions:** An examination of hearing impairment must be conducted by a state-licensed audiologist and must include a controlled speech discrimination test (specifically, the Maryland CNC recording) and a puretone audiometry test in a sound isolated booth that meets American National Standards Institute standards (ANSI S3.1.1999 [R2004]) for ambient noise. Measurements will be reported at the frequencies of 500, 1000, 2000, 3000, and 4000 Hz.

The examination will include the following tests: Puretone audiometry by air conduction at 250, 500, 1000, 2000, 3000, 4000, 6000 Hz and 8000 Hz, and by bone conduction at 250, 500, 1000, 2000, 3000, and 4000 Hz, spondee thresholds, speech discrimination using the recorded Maryland CNC Test, tympanometry and acoustic reflex tests (ipsilateral and contralateral), and, when necessary, Stenger tests. Bone conduction thresholds are measured when the air conduction thresholds are poorer than 15 dB HL. A modified Hughson-Westlake procedure will be used with appropriate masking. A Stenger must be administered whenever puretone air conduction thresholds at 500, 1000, 2000, 3000, and 4000 Hz differ by 20 dB or more between the two ears.

Maximum speech discrimination will be reported with the 50 word VA approved recording of the Maryland CNC test. The starting presentation level will be 40 dB re SRT. If necessary, the starting level will be adjusted upward to obtain a level at least 5 dB above the threshold at 2000 Hz, if not above the patient's tolerance level.

The examination will be conducted without the use of hearing aids. Both ears must be examined for hearing impairment even if hearing loss in only one ear is at issue.

When speech discrimination is 92% or less, a performance intensity function must be obtained.

A comprehensive audiological evaluation should include evaluation results for puretone thresholds by air and bone conduction (500-8000 Hz), speech reception thresholds (SRT), speech discrimination scores, and acoustic immittance with acoustic reflexes (ipsilateral and contralateral reflexes). Tests for non-organicity must be performed when indicated.

**1. OBJECTIVE FINDINGS****A. PURETONE THRESHOLDS IN DECIBELS (AIR CONDUCTION):**

Instructions: Measure and record puretone threshold values in decibels at the indicated frequencies (air conduction). Report the decibel (dB) value, which ranges from -10 dB to 105 dB, for each of the frequencies. Add a plus behind the decibel value when a maximum value has been reached with a failure of response from the Veteran. In those circumstances where the average includes a failure of response at either the maximum allowable limit (105 dB) or the maximum limits of the audiometer, use this maximum decibel value of the failure of response in the puretone threshold average calculation.

If the Veteran could not be tested (CNT), enter CNT and state the reason why the Veteran could not be tested. Clearly inaccurate, invalid or unreliable test results should not be reported.

The puretone threshold at 500 Hz is not used in calculating the puretone threshold average for evaluation purposes but is used in determining whether or not for VA purposes, hearing impairment reaches the level of a disability. The puretone threshold average requires the decibel levels of each of the required frequencies (1000 Hz, 2000 Hz, 3000 Hz, and 4000 Hz) be recorded for the test to be valid for determination of a hearing impairment.

**RIGHT EAR**

	A	B	C	D	E	F	G	
	500 Hz*	1000 Hz*	2000 Hz*	3000 Hz*	4000 Hz*	6000 Hz*	8000 Hz*	Avg Hz (B-E)**
	15	15	25	25	25	20	25	22.5

**LEFT EAR**

	A	B	C	D	E	F	G	
	500 Hz*	1000 Hz*	2000 Hz*	3000 Hz*	4000 Hz*	6000 Hz*	8000 Hz*	Avg Hz (B-E)**
	20	15	25	25	30	20	35	23.75

\*The puretone threshold at 500 Hz is not used in determining the evaluation but is used in determining whether or not a ratable hearing loss exists.

\*\*The average of B, C, D, and E.

\*\*\*CNT - Could Not Test

**B. WERE THERE ONE OR MORE FREQUENCY(IES) THAT COULD NOT BE TESTED?**

☐ YES ☒ NO *If yes, enter CNT in the box for frequency(ies) that could not be tested, and explain why testing could not be done:*

**C. VALIDITY OF PURETONE TEST RESULTS:**

- ☒ Test results are valid for rating purposes.
- ☐ Test results are not valid for rating purposes (not indicative of organic hearing loss).
- If invalid, provide reason:

**D. SPEECH DISCRIMINATION SCORE (MARYLAND CNC WORD LIST)**

Instructions on pausing: Examiners should pause when necessary during speech discrimination tests, in order to give the Veteran sufficient time to respond. This will ensure that the test results are based on actual hearing loss rather than on the effects of other problems that might slow a Veteran's response. There are a variety of problems that might require pausing, for example, the presence of cognitive impairment. It is up to the examiner to determine when to use pausing and the length of the pauses.

RIGHT EAR	100	%
LEFT EAR	100	%

**E. APPROPRIATENESS OF USE OF WORD RECOGNITION SCORE (MARYLAND CNC WORD LIST):**

RIGHT EAR:

IS WORD DISCRIMINATION SCORE AVAILABLE?

☒ YES ☐ NO☒ Use of speech discrimination score is appropriate for this Veteran.☐ The use of the speech discrimination score is not appropriate for this Veteran because of language difficulties, cognitive problems, inconsistent speech discrimination scores, etc., that make combined use of puretone average and speech discrimination scores inappropriate.

LEFT EAR:

IS WORD DISCRIMINATION SCORE AVAILABLE?

☒ YES ☐ NO☒ Use of speech discrimination score is appropriate for this Veteran.☐ The use of the speech discrimination score is not appropriate for this Veteran because of language difficulties, cognitive problems, inconsistent speech discrimination scores, etc., that make combined use of puretone average and speech discrimination scores inappropriate.**F. AUDIOLOGIC FINDINGS**

Summary of Immittance (Tympanometry) Findings:

	RIGHT EAR		LEFT EAR	
ACOUSTIC IMMITTANCE	<input checked="" type="checkbox"/> Normal	<input type="checkbox"/> Abnormal	<input checked="" type="checkbox"/> Normal	<input type="checkbox"/> Abnormal
IPSILATERAL ACOUSTIC REFLEXES	<input checked="" type="checkbox"/> Normal	<input type="checkbox"/> Abnormal	<input checked="" type="checkbox"/> Normal	<input type="checkbox"/> Abnormal
CONTRALATERAL ACOUSTIC REFLEXES	<input checked="" type="checkbox"/> Normal	<input type="checkbox"/> Abnormal	<input checked="" type="checkbox"/> Normal	<input type="checkbox"/> Abnormal
UNABLE TO INTERPRET REFLEXES DUE TO ARTIFACT	<input type="checkbox"/>		<input type="checkbox"/>	
UNABLE TO OBTAIN / MAINTAIN SEAL	<input type="checkbox"/>		<input type="checkbox"/>	

**2. DIAGNOSIS**RIGHT EAR

<input checked="" type="checkbox"/> Normal hearing	ICD CODE:
<input type="checkbox"/> Conductive hearing loss	ICD CODE:
<input type="checkbox"/> Mixed hearing loss	ICD CODE:
<input type="checkbox"/> Sensorineural hearing loss (in frequency range of 500-4000 Hz)*	ICD CODE:
<input type="checkbox"/> Sensorineural hearing loss (in frequency range of 6000 Hz or higher frequencies)**	ICD CODE:
<input type="checkbox"/> Significant changes in hearing thresholds in service***	ICD CODE:

LEFT EAR

<input type="checkbox"/> Normal hearing	ICD CODE:
<input type="checkbox"/> Conductive hearing loss	ICD CODE:
<input type="checkbox"/> Mixed hearing loss	ICD CODE:
<input checked="" type="checkbox"/> Sensorineural hearing loss (in frequency range of 500-4000 Hz)*	ICD CODE: H90
<input checked="" type="checkbox"/> Sensorineural hearing loss (in frequency range of 6000 Hz or higher frequencies)**	ICD CODE: H90
<input type="checkbox"/> Significant changes in hearing thresholds in service***	

**NOTES:**

\*The Veteran may have hearing loss at a level that is not considered to be a disability for VA purposes. This can occur when the auditory thresholds are greater than 25 dB at one or more frequencies in the 500-4000 Hz range.

\*\* The Veteran may have impaired hearing, but it does not meet the criteria to be considered a disability for VA purposes. For VA purposes, the diagnosis of hearing impairment is based upon testing at frequency ranges of 500, 1000, 2000, 3000, and 4000 Hz. If there is no HL in the 500-4000 Hz range, but there is HL above 4000 Hz, check this box.

\*\*\*The Veteran may have a significant change in hearing threshold in service, but it does not meet the criteria to be considered a disability for VA purposes. (A significant change in hearing threshold may indicate noise exposure or acoustic trauma.)

**3. ETIOLOGY**☐ ETIOLOGY OPINION NOT INDICATED AS: ☐ SERVICE CONNECTED CONDITION ☐ VBA DID NOT REQUEST ETIOLOGYRIGHT EAR

WAS THERE A PERMANENT POSITIVE THRESHOLD SHIFT (WORSE THAN REFERENCE THRESHOLD) GREATER THAN NORMAL MEASUREMENT VARIABILITY AT ANY FREQUENCY BETWEEN 500 AND 6000 HZ FOR THE RIGHT EAR?

☐ YES ☒ NO

OPINION PROVIDED FOR THE RIGHT EAR:

☐ YES ☒ NO

### 3. ETIOLOGY (continued)

#### RIGHT EAR (continued)

IF PRESENT, IS THE VETERAN'S RIGHT EAR HEARING LOSS AT LEAST AS NOT (50% PROBABILITY OR GREATER) CAUSED BY OR A RESULT OF AN EVENT IN MILITARY SERVICE?

☐ YES

☐ NO

☐ CANNOT DETERMINE A MEDICAL OPINION REGARDING THE ETIOLOGY OF THE VETERAN'S RIGHT EAR HEARING LOSS WITHOUT RESORTING TO SPECULATION:

RATIONALE (Provide rationale for either a yes, no answer or speculation reason):

DID HEARING LOSS EXIST PRIOR TO SERVICE?

☐ YES ☐ NO

IF YES, WAS THE PRE-EXISTING HEARING LOSS AGGRAVATED BEYOND NORMAL PROGRESSION IN MILITARY SERVICE?

☐ YES ☐ NO

PROVIDE RATIONALE FOR BOTH YES OR NO:

#### LEFT EAR

WAS THERE A PERMANENT POSITIVE THRESHOLD SHIFT (WORSE THAN REFERENCE THRESHOLD) GREATER THAN NORMAL MEASUREMENT VARIABILITY AT ANY FREQUENCY BETWEEN 500 AND 6000 HZ FOR THE LEFT EAR?

☐ YES ☒ NO

OPINION PROVIDED FOR THE LEFT EAR:

☒ YES ☐ NO

IF PRESENT, IS THE VETERAN'S LEFT EAR HEARING LOSS AT LEAST AS NOT (50% PROBABILITY OR GREATER) CAUSED BY OR A RESULT OF AN EVENT IN MILITARY SERVICE?

☒ YES

☐ NO

☐ CANNOT DETERMINE A MEDICAL OPINION REGARDING THE ETIOLOGY OF THE VETERAN'S LEFT EAR HEARING LOSS WITHOUT RESORTING TO SPECULATION:

RATIONALE (Provide rationale for either a yes, no answer or speculation reason):

Record showing MOS of Special Forces which had a High probability for hazardous noise exposure. 1963 and 1975 both showed normal hearing sensitivity; current exam shows clinical hearing loss.

DID HEARING LOSS EXIST PRIOR TO SERVICE?

☐ YES ☒ NO

IF YES, WAS THE PRE-EXISTING HEARING LOSS AGGRAVATED BEYOND NORMAL PROGRESSION IN MILITARY SERVICE?

☐ YES ☐ NO

PROVIDE RATIONALE FOR BOTH YES OR NO:

### 4. FUNCTIONAL IMPACT OF HEARING LOSS

NOTE: Ask the Veteran to describe in his or her own words the effects of disability (i.e., the current complaint of hearing loss on occupational functioning and daily activities). Document the Veteran's response without opining on the relationship between the functional effects and the level of impairment (audiogram) or otherwise characterizing the response. Do not use handicap scales.

DOES THE VETERAN'S HEARING LOSS IMPACT ORDINARY CONDITIONS OF DAILY LIFE, INCLUDING ABILITY TO WORK?

☒ YES ☐ NO

IF YES, DESCRIBE IMPACT IN THE VETERAN'S OWN WORDS:

Difficulty understanding those around me. Required to stand close to understand.  
Still on active duty as special forces.



**5. REMARKS, IF ANY, PERTAINING TO HEARING LOSS:**

Pre-military: None During-military: M9 handgun, M16 rifle, M4 Rifle, M249 Rifle, hand grenade training, right handed shooter. 30+ years as a active duty special forces. aircraft.

The examinee reports serving in the Army. The examinee indicates they served a total of 40+ year(s). The period(s) of service were from: 11/17/1965 to 7/31/2020. The examinee was in service during Vietnam War, Gulf War, Afghan War and Iraq War. The examinee reports that they participated in combat activity.

For the claimant's claimed condition of hearing loss, left please refer to the diagnosis

**SECTION 2: TINNITUS****1. MEDICAL HISTORY**

DOES THE VETERAN REPORT RECURRENT TINNITUS?

☒ YES ☐ NO

DATE AND CIRCUMSTANCES OF ONSET OF TINNITUS:

Approximately 20 years ago after a night of flying on a C130 I noticed a ringing in my ears while trying to go to sleep. The reported tinnitus is constant. The side(s) affected:

Both

**2. ETIOLOGY OF TINNITUS**

SELECT ANSWER BELOW AND PROVIDE RATIONALE WHERE REQUESTED:

- ☐ ETIOLOGY OPINION NOT INDICATED AS: ☐ SERVICE CONNECTED CONDITION ☐ VBA DID NOT REQUEST ETIOLOGY
- ☐ THE VETERAN HAS A DIAGNOSIS OF CLINICAL HEARING LOSS, AND HIS OR HER TINNITUS IS AT LEAST AS LIKELY AS NOT (50% PROBABILITY OR GREATER) A SYMPTOM ASSOCIATED WITH THE HEARING LOSS, AS TINNITUS IS KNOWN TO BE A SYMPTOM ASSOCIATED WITH HEARING LOSS.
- ☐ LESS LIKELY THAN NOT (LESS THAN 50% PROBABILITY) A SYMPTOM ASSOCIATED WITH THE VETERAN'S HEARING LOSS

RATIONALE:

- ☒ AT LEAST AS LIKELY AS NOT (50% PROBABILITY OR GREATER) CAUSED BY OR A RESULT OF MILITARY NOISE EXPOSURE

RATIONALE:

Veteran clearly reports it secondary to a military event

- ☐ AT LEAST AS LIKELY AS NOT (50% PROBABILITY OR GREATER) DUE TO A KNOWN ETIOLOGY (*such as traumatic brain injury*)

RATIONALE:

- ☐ LESS LIKELY THAN NOT (LESS THAN 50% PROBABILITY) CAUSED BY OR A RESULT OF MILITARY NOISE EXPOSURE

RATIONALE:

- ☐ CANNOT PROVIDE A MEDICAL OPINION REGARDING THE ETIOLOGY OF THE VETERAN'S TINNITUS WITHOUT RESORTING TO SPECULATION

REASON SPECULATION REQUIRED:

### 3. FUNCTIONAL IMPACT OF TINNITUS

NOTE: Ask the Veteran to describe in his or her own words the effects of disability (i.e., the current complaint on occupational functioning and daily activities). Document the Veteran's response without opining on the relationship between the functional effects and the level of impairment (audiogram) or otherwise characterizing the response. Do not use handicap scales.

DOES THE VETERAN'S TINNITUS IMPACT ORDINARY CONDITIONS OF DAILY LIFE, INCLUDING ABILITY TO WORK?

☒ YES ☐ NO

IF YES, DESCRIBE IMPACT IN THE VETERAN'S OWN WORDS

Ringling makes it difficult to focus for sustained periods.

### 4. REMARKS, IF ANY, PERTAINING TO TINNITUS

### SECTION 3: PHYSICIAN'S CERTIFICATION AND SIGNATURE

**CERTIFICATION** - To the best of my knowledge, the information contained herein is accurate, complete and current.

3A. AUDIOLOGIST/PHYSICIAN SIGNATURE & TITLE

3B. AUDIOLOGIST/PHYSICIAN PRINTED NAME

Richard Smith M.Au.D Audiology

3C. DATE SIGNED

3D. AUDIOLOGIST/PHYSICIAN PHONE AND FAX NUMBER

619-400-1234

3E. NATIONAL PROVIDER IDENTIFIER (NPI) NUMBER

3F. AUDIOLOGIST/PHYSICIAN ADDRESS

NPI: 12566666 Lic#:A#697 AR

8810 Rio San Diego Drive, San Diego, CA 92108

**NOTE** - VA may request additional medical information, including additional examinations, if necessary to complete VA's review of the veteran's application.

**IMPORTANT** - Audiologist/Physician please fax the completed form to \_\_\_\_\_

(VA Regional Office FAX No.)

**NOTE** - A list of VA Regional Office FAX Numbers can be found at [www.benefits.va.gov/disabilityexams](http://www.benefits.va.gov/disabilityexams) or obtained by calling 1-800-827-1000.

**PRIVACY ACT NOTICE:** VA will not disclose information collected on this form to any source other than what has been authorized under the Privacy Act of 1974 or Title 38, Code of Federal Regulations 1.576 for routine uses (i.e., civil or criminal law enforcement, congressional communications, epidemiological or research studies, the collection of money owed to the United States, litigation in which the United States is a party or has an interest, the administration of VA programs and delivery of VA benefits, verification of identity and status, and personnel administration) as identified in the VA system of records, 58/VA21/22/28, Compensation, Pension, Education and Vocational Rehabilitation and Employment Records - VA, published in the Federal Register. Your obligation to respond is voluntary. VA uses your SSN to identify your claim file. Providing your SSN will help ensure that your records are properly associated with your claim file. Giving us your SSN account information is voluntary. Refusal to provide your SSN by itself will not result in the denial of benefits. VA will not deny an individual benefits for refusing to provide his or her SSN unless the disclosure of the SSN is required by a Federal Statute of law in effect prior to January 1, 1975, and still in effect. The requested information is considered relevant and necessary to determine maximum benefits under the law. The responses you submit are considered confidential (38 U.S.C. 5701). Information submitted is subject to verification through computer matching programs with other agencies.

**RESPONDENT BURDEN:** We need this information to determine entitlement to benefits (38 U.S.C. 501). Title 38, United States Code, allows us to ask for this information. We estimate that you will need an average of 30 minutes to review the instructions, find the information, and complete the form. VA cannot conduct or sponsor a collection of information unless a valid OMB control number is displayed. You are not required to respond to a collection of information if this number is not displayed. Valid OMB control numbers can be located on the OMB Internet Page at [www.reginfo.gov/public/do/PRAMain](http://www.reginfo.gov/public/do/PRAMain). If desired, you can call 1-800-827-1000 to get information on where to send comments or suggestions about this form.

For Internal VA Use

Hearing Loss and Tinnitus Disability Benefits Questionnaire

Updated on: March 31, 2020 ~v20\_1

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