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PROJECT RANCH HAND II

EXAMINER'S HANDBOOK

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A. General Instructions

Project RANCH HAND II is a multiyear effort to determine whether or not C-123 aircrew members who were engaged in the aerial spraying of herbicides in Vietnam have developed significant adverse health effects from that exposure. Detailed surveys of the world's literature have been used in designing the history questionnaires, physical examination protocol, and laboratory procedures.

This phase of Project RANCH HAND II involves a cross sectional study of the subject's health at the time of examination. It is important that examiners remain unaware of the subject's status as a RANCH HAND participant or as a control subject. The physician examiner is tasked to examine and objectively record his findings. The examining physician is not, and cannot be expected to arrive at any definitive diagnosis as the full history and laboratory results are not available to him. The compilation and analysis of data will be performed by the study investigators at Brooks Air Force Base, Texas. They will notify the subject and the physician of his choice of the results of the examination.

The physicians performing examinations for Project RANCH HAND II should be aware that the report of examination will become a permanent record. This report will be referred to not only in the near future as the cross sectional study is analyzed, but at the time of the next review of the subject in the follow-up phases of Project RANCH HAND. These examinations will define the health status of the subjects at a point in time, and to establish the presence of physical findings, if any exist. After statistical review of the study groups, these findings may permit definition of a chronic effect due to exposure. An inaccurate examination may lead to falacious study results in two ways: a presumed syndrome may be defined which does not in fact exist, or a syndrome which in fact exists may not be defined with enough validity to warrant further actions.

The examining physician is responsible for recording a complete and detailed report of the physical examination. In this role, the examining physician is tasked with collecting evidence of the presence or absence of physical signs of abnormality only. Formulation of impressions is not requested nor desired. All items on the physical examination report form must be completed. It is imperative that the physician make such additional remarks as may be required to adequately describe existing physical and mental impair-The examining physician must avoid an expression of opinion ments. regarding the interpretation of any findings particularly with regards to possible etiologies. If, during the examination, the physician discovers evidence of acute serious illness requiring immediate treatment, the normal emergency or urgent care procedures of the medical facility would apply. If during the examination, the examining physician finds evidence of present illness requiring further medical attention, he should so state to the subject and offer to forward or have forwarded pertinent information to the subject's physician. A clear record of any such advice and treatment should be recorded. The ultimate value of the RANCH HAND II Study will lie in complete, accurate and, whenever possible, quantitative data permitting the most stringent and powerful statistical analysis. For that reason, the physical examination protocol requires exact measurements in many instances, and the use of defined meanings of semiquantitative indicators in other places.

B. Conduct of the Examination

(1) On arrival at the examining facility the subject should be briefed on the appointments which have been arranged, their times, and locations.

(2) Collation and forwarding of examination results

A checklist for the mailing of data will be provided. It should be retained by the office primarily responsible (OPR) for the Project RANCH HAND II examination and used to ascertain that all necessary items have been completed and received, or have been directly forwarded by the section performing the examinations. When the OPR for the examinations is ready to forward all materials, the checklist for mailing should be endorsed with the date of mailing as a letter of transmittal and included in the package of material to be mailed to USAFSAM/EK, Brooks AFB TX 78235.

(3) Forms for individual examinations and procedures

The blank forms included for various examinations and procedures may be carried by the patient so as to be available to the examiner or to the laboratory, or to the department of radiology, as the patient reports for his examinations in those functions. The forms pertaining to the specific function may be withdrawn from the patient's examination package and later returned to the office of primary responsibility.

	C. Londuct of the	Examination	
SECTION	PHYSICAL EXAMINATION	<u></u>	SUBJECT NUMBER
e. 7 Well-nourished (3) 7 Older as h. Appearance of illr	/ Mesomorph c. / 7 d f. / 7 Obese g. App stated age. mess or distress / 7 Ye / 7 Normal / 7 Abnorm	earance (1) <u>7</u> Younge es <u>7</u> No	er (2) <u>/</u> Same
2. HEIGHT CM	WEIGHT (Undrogged) Kg	SITTING BLOOD PRESSURE R	IGHT ARM AT HEART LEVEL
2. PULSE RATE REGUL		SYSTOLIC Describe any irregularities	
a. Irregular / b. Irregularly irregu c. VPBs per minute	lar <u>7</u>	· • •	
4 EYE GROUNDS []NORM/ [7] A-V nicking [7] † light reflex [7] Arteriolar spasm	Hemorrhage ^g pages T Exudates	7 Papilledema	rhages, exudates,
	ASNORMAL Des tact /7 Yes /7 No		gmentation
7. NECK (Repocially thyroid gland) Thyroid gland palpabl Enlarge Nodule Tenderness	e / / Parotid g d / / R s / R	ORMAL Describe any abnor gland enlargement 7 L	mality.
a. THORAX AND LUNGS // Asymmetrical expa // Hyperresonance // Dullness		AL Describe any abnormality, Circumference at n Expiration Inspiration	
9. HEART [] NORMAL Displacement of apica Heart sounds normal (Continued in Item 18 10. ABDOMEN [] Heptomegaly [] Heptomegaly [] Cm Liver Span [] Splenomegaly	l impulse / No / Y / Yes / No / Si on Reverse) #ORMAL D	éscribe aby abnomality with sp pleen and liver. Record wa	edal attention to the ist measurement ed form.
11. EXTREMITIES []]NORM // Absence, specify: // Edema // Pitting // 1 // Loss of hair on to // R // L	AAL ABNORMAL Der Non-pitting 7 V	scribe any edema or signs of ve Lubbing of nails aricosities	

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SECTION '	HYSICAL EX	AMINATIO	H (Contin	ied)	······································
12. PERIPHERAL PULSES	NORMAL	DIMIN.	ABSEN T	COMMENTS	······································
RADIAL			†		
FENORAL				•	
POPLITEAL					
DORSALIS PEDIS					
POSTERIOR TIBIAL			 		
13. SKIN			····· •		
Comedones Acneiform lesions Acneiform scars Depigmentation Inclusion cysts		Hyper Jaund Spide Palma	att pigmen lice r angi r eryt	omata <u>7</u> Petechiae Ecchymoses	ds Light
14. MUSCULOSKELETAL Muscle - Specify: // Weakness // Tenderness // Abnormal Cons // Atrophy	4	NORM.	ine 7 Scol 7 Kyph 7 Tend Le 7 Decr	ABNORMAL Losis Dosis erness, Zel eased range /7 Pelvic motion /7 Straig	
15. GENITOURINARY - RECT // Inguinal hernia // Testes Absent E // R // L	∠7R 4 nlarged ∠7 ∠7	Atropi		AL / ABNORMAL Varicocele Epididymis Scotal mass - Specify cm dia Positive	Hemorrhoids Prostatic Enlargement Rectal mass Negative
OCCIPITAL, SUPR	ACLAVICUI	LAR, Ander <u>(</u>		PRMAL / ABNORMAL - SPEC: , EPITRACHLEAR, INGUINAL, 7 Hard / Fixed	FEMORAL
18. HEART AND OTHER OF	BSERVATIC				
(Continued from Item 9) Murmur // No // Yes		7 Ao	Z F	u _7 Apex _7 54	
· · · ·	Sys /				
DATE OF EXAMINATION				ED NAME OF EXAMINING PHYSICIAN	RETURN FORM
MONTHDAY	YEAR	1		• · · · · · · · · · · · · · · · · · · ·	TO:
AMINING FACILITY		SIGNATU	AE		USAFSAM/ ES BROOKS AFB
		ł			TX 78235

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CLINICAL RECORD NEUROLOGICAL EXAMINATION HEAD AND NECK - Normal to Palpations/Inspection / 7Y / 7N Specify Scar / 7 Asymmetry /7 Depression /7 Carotid Bruit _7No _7R _7L Neck Range of Motion /7 Normal or Decreased to /7 Left /7 Right /7 Forward /7 Backward TRUNK MOTOR SYSTEM - Handedness Right /7 Left /7 Gait [7] Normal or /7 Broad Based /7 Ataxic /7Small Stepped /7Other-Specify Associated Movements / Arm Swing / [Norma] or Abnormal / 7R / 7L Muscle Status (strength, tone, volume, tenderness, fibrillations) Bulk /7 Normal /7 Abnormal Tone Upper Extremities / Normal or / 7Increased / 7Decreased /7Right /7Left Lower Extremities / Normal or / Increased / Decreased /7Right /7Left Strength - Distal wrist extensors / 7Normal / 7Decreased Ankle/Toe Dors/Flexors / Normal / Decreased / R / L Proximal Deltoids / 7Normal / 7Decreased / 7R / 7L Hip Flexors /7Normal /7Decreased /7R /7L Abnormal Movements (tremors, tics, choreas, etc.) Fasiculations / 7No / Yes (1-4+) Tenderness / 7No / 7Yes (1-4+) Tremor / 7No / 7Yes - Specify Upper Extremity / R / Resting / Essential / Intention Lower Extremity / R / L / Other Coordination (a) Equilibratory - Eyes Open Eyes Closed - Romberg ____Positive (Abnormal) ____Negative (Normal) **Right Foot** Left Foot If indicated, check / Normal / Abnormal / R / R Rapidly alternative movements / 7Normal / 7Abnormal / 7R / 7L / 7Both Skilled Acts (a) Praxis (b) Handwriting. If indicated, []Normal []Abnormal (c) Speech (articulation, aphasia, agnosia) Grossly / 7Normal /7Abnormal - Specify Dysarthria /7 Aphasia /7

Reflexes (0-absent; 1-sluggish; 2-active; 3-very active; 4-transient clonus; 5-sustained clonus) R R L Abnormal R L **Other** R Deep L Deep L Babinski Patellar Biceps Triceps Achilles Remarks MENINGEAL IRRITATION Spurling Maneuver of Neck []Normal []Abnormal /7R /7L /─7Both Straight Leg Raising ____Normal ____Abnormal ____R ___L ___Both NERVE STATUS (tenderness, tumors, etc.) SENSORY SYSTEM (tactile, pain, vibration, position. If positive sensory signs are present, summarize below and indicate details on Anatomical Figure, Std. Form 531) Light Touch / 7Normal / 7Abnormal (Map on Anatomical Figure) Pin Prick / 7Normal / 7Abnormal Vibration (at ankle, 128 hz tuning fork): ___Normal ___Abnormal ___R ___L ___Both Position (Great toe): /7Normal /7Abnormal /7R /1L //Both CRANIAL NERVES I R Smell / Present / Absent L Smell / Present / Absent II Fundus R Normal [7] Abnormal [7] Disk Pallor/atrophy [] Exudate [] Papilledema [] Hemorrhage Fundus L Normal / Abnormal / Disk pallor/atrophy //Exudate / Papilledema / Hemorrhage Fields (to confrontation) Right /___Normal /___Abnormal Left /___Normal /___Abnormal III Normal / / / Abnormal - Specify Pupils-Size (mm) Equal [7] Unequal [7] Difference mm_____ Shape, position Round [7] Other [7] [7]R [7]L Light, Reaction Normal [7] Abnormal [7] [7]R [7]L I٧ Ϋİ Position of Eyeballs L Movements R Nystagmus Rotary [7] Horizontal [7] Vertical [7] (Draw position)

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Ptosis R/7 V Motor R Clench Jaw - Symmetric [7] Deviated [7] R[7] L[7] Sensory R Normal 7 Abnormal 7 V17 L Normal 7 Abnormal 7 V17 V_2 V_3 V_2 V_3 Corneal Reflex R L VII Motor R Normal smile / Yes / No Palpebral Fissure / Yes / No L Normal smile / Yes / No Palpebral Fissure / Yes / No IX Palate and Uvula \times Movement Normal \square Deviation to $\square R$ $\square L$ Palatal Reflex R /7Normal /7Abnormal L [7Norma] [7Abnorma] XII Tongue-Protruded-Central / 7 R / 7 L / 7 Atrophy ___No ___Yes MENTAL STATUS (alert, clear, cooperative, etc.) Gross abnormalities: ///No / 7Yes - Specify

> SUMMARY OF POSITIVE FINDINGS Objective

Subjective

Diagnostic Impression

Date

Signature

D. Special Procedures

(1) Nerve Conduction Velocities

(a) These studies have been determined to be an important parameter in long-term follow-up studies of persons thought to have been exposed to Herbicide Orange Components.

(b) The Nerve Conduction Velocities should be performed by a physician or by a specialty qualified technician under the supervision of a physician trained in neurophysiological methods.

- (c) <u>Specific NCVs</u> (See form included in F. Below)
 - (1) Ulnar Nerve (one side only)
 - (a) motor (above elbow, below elbow)
 - (b) values recorded
 - (i) distal latency
 - (ii) NCV
 - (2) <u>Peroneal Nerve</u> (one side only)
 - (<u>a</u>) motor
 - (b) values recorded
 - (i) distal latency
 - (ii) NCV
 - (<u>3</u>) <u>Sural Nerve</u> (one side only)
 - (a) sensory: orthodromic
 - (b) values recorded: NCV
- (d) <u>Methods</u>

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PERONEAL NERVE

(1) Active electrode is placed over the extensor digitorum brevis and reference over the little toe. Stimulating electrodes are placed over anterior distal leg 8 cm proximal to active electrode. Proximal site is distal to head of fibula. If entrapment is suspected at fibular head use a stimulation site of 12-18 cm more proximal to the fibular head.

Anomalous innervation to the extensor digitorum brevis occurs in 1/5 patients (at least partially). Identified by inability to evoke a muscle action potential when stimulating at anterior ankle or a different shape (smaller) potential when stimulating here. This accessory nerve causes posterior to lateral malleolus so cathode should be placed here.

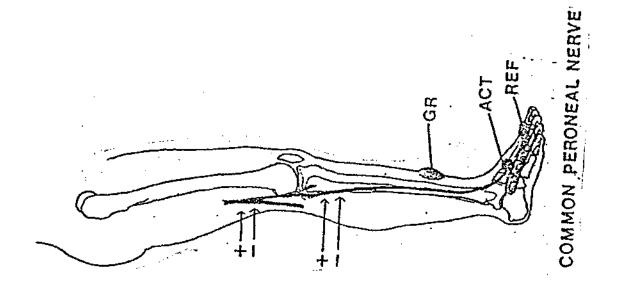
NORMAL VALUES

 49.9 ± 5.9 M/sec Distal latency: $4.5 \pm .8$ ms

Proximal latencies have been determined for use in below the knee amputees, and neuromuscular diseases where extensor digitorum brevis action potential cannot be elicited. Active electrode is placed 1/2 way down leg over middle of dorsiflexor muscle group and stimulation at fibular head.

NORMAL VALUES

5.5 - 7.2 ms (N = 217)



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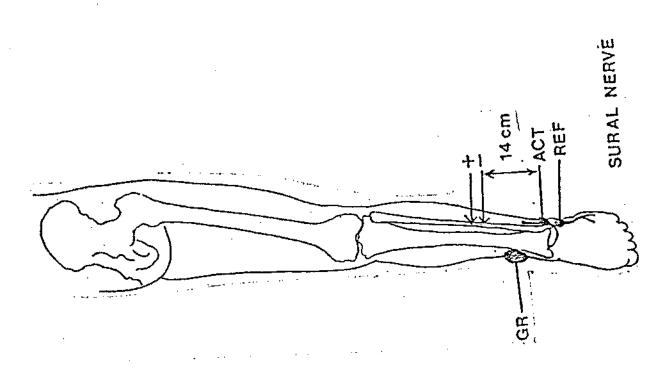
9

SURAL NERVE

 $(\underline{2})$ Active and recording electrodes are placed under lateral malleolus on lateral aspect of ankle. Sural nerve is stimulated as it pierces the gastrocnemius fascia just lateral to the midline of posterior distal calf, 10-18 cm proximal to active electrode. If leg is cold - a clue is prolonged latency of peroneal nerve - determine temperature. Subtract .1 ms (latency of activation) from the observed latency and divide into the distance.

NORMAL VALUES (after LaFratta)

Age	(To Peak)
20-29	44 ± 2.5 M/sec
30-39	38.80 ± 3.3 M/sec
40-49	36.70 ± 3.7 M/sec
50-59	37.20 ± 3.0 M/sec
60 α over	35.00 ± 3.8 M/sec



ULNAR NERVE

MOTOR CONDUCTION

(3) Active electrode is placed over center of abductor digiti quinti; reference over proximal phalanx fifth digit. Stimulation (cathode) just radial to tendon of flexor carpi ulnaris 8 cm proximal to active electrode. Proximal site of stimulation should be just below ulnar groove and 18 cm proximal to ulnar groove on medial aspect of humerus.

<u>N.B.</u>: Elbow should be flexed to 70 degrees during procedure of stimulation and measurement to make more precise the actual length of ulnar nerve. More proximal stimulation sites include supraclavicular and C-8 root (see median nerve).

SENSORY CONDUCTION

Antidromic - ring electrodes over fifth digit separated by 4 cm. N.B. motor artifact may be interfering. Stimulate 14 cm proximal to active electrode at same site as motor stimulation.

Orthodromic - reverse stimulation and recording electrodes. More proximal sites of stimulation may also be done.

NORMAL VALUES

 57 ± 4.7 M/sec - motor forearm segment 62.7 \pm 5.5 M/sec - motor across elbow segment 56.7 \pm 4.2 M/sec - sensory orthodromic (to peak) 54.9 \pm 3.9 M/sec - sensory antidromic (to peak)

Distal Latency:

Motor: $3.7 \pm .3$ Sensory: $3.0 \pm .25$ Antidromic (peak) $3.0 \pm .25$ Orthodromic (peak)

Muscle AP 8-20 mV

Sensory AP 15-50 mV

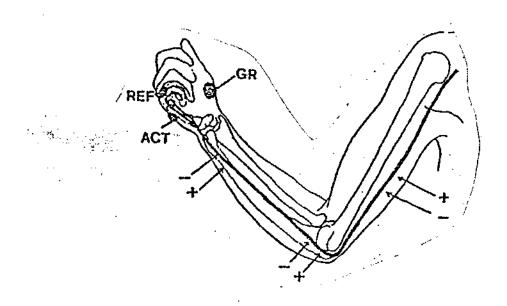
ADDENDUM

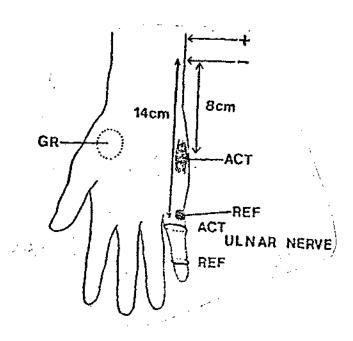
For deep branch surface recording electrode should be over adductor pollicus (i.e. just medial to thenar eminence on palmar surface of web space). Additional latency is .5 ms.

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(e) Disposition

Forward the recorded results on the form attached to the examination package to the examining physician.

(2) Psychological Battery

(a) <u>General</u>

(1) This battery yields objective numerical data, and is well-standardized and clinically validated. The individual tests were chosen to insure an adequate analysis of one of the major alleged manifestations of herbicide toxicity. Each test either validates one of the other tests, or is considered to be a "definitive" test for analysis of a suspected psycho/neuropathic effect.

(2) Compared to the general civilian population, characteristic response tendencies are observed on the MMPI and Cornell Index among active duty aircrewmen being evaluated in an aeromedical setting. It is also important to consider the effect that pending retirement has exerted on the reporting of medical history and symptomatology. This may also alter responses to psychological testing.

(3) The battery requires approximately 5-1/2 to 6-3/4 hours to administer, depending on the speed of the examinee. An additional 1 to 2 hours of scoring and other clerical tasks will be required. Since test debriefing to clarify unusual performances, response biases, etc., is a crucial part of the psychologic evaluation, it is recommended that testing begin and be completed as early as possible during each examinee's stay at his respective evaluative facility.

(b) <u>Specific Tests</u>

(<u>1</u>) <u>Wechsler Adult Intelligence Scale</u> (<u>WAIS</u>): 60-75 minute individually-administered collection of verbal and nonverbal intellectual measures; also useful for clinical inferences when combined with the neuropsychological battery below.

(2) <u>Reading subtest of the Wide Range</u> <u>Achievement Test (WRAT)</u>: 10-minute individually-administered measure of word recognition ability. Important so as to rule-out reading inefficiency should response to personality instruments below be of questionable validity (e.g., high F Scale on MMPI).

(3) <u>Halstead-Reitan Neuropsychological Test</u> <u>Battery</u>: 150-180 minute individually-administered collection of brain behavior relationship measures for establishing the functional integrity of the cerebral hemispheres. The battery must include the following subtests: Category, Tactual performance, Speech-Sounds,

Seashore Rhythm, Finger Tapping, Trail Making, and Grip Strengths. The Aphasia Screening and Sensory-Perceptual Exams are considered optional in view of their redundancy with the clinical neurologic exam included in this project. Individualized test debriefing is conducted to clarify test performances in the WAIS and Neuropsychological Battery.

(4) <u>Three subtests of the Wechsler Memory</u> <u>Scale I (WMS I)</u>: 30-minute individually-administered measures of immediate and delayed recall of verbal and visual materials. The Logical Memory, Associate Learning and Visual Reproduction subtests are to be administered in the standard, immediate-recall fashion initially. After 30 minutes has elapsed, the examinee is asked, without prior alerting, to recall as much as he can about the Logical Memory and Visual Reproduction subtest stimuli. Standard scoring is used for both test-retest administrations.

(5) <u>Cornell Index (CI)</u>: 10-15 minute selfadministered and standardized neuropsychiatric symptom and complain inventory, including items involving asthenia, depression, anxiety, fatigue, and GI symptoms in lay language. Endorsement of items are to be explored and clarified in test-debriefing.

(6) <u>Minnesota Multiphasic Personality</u> <u>Inventory (MMPI)</u>: 60 to 90 minute self administered clinical psychiatric screening instrument; also capable of estimating response biases (e.g., "fake good," or "fake bad"). The shortened version of Form R (i.e., items 1 to 399) may be substituted for the 566-item Long Form. Standard scoring and Minnesota norms are to be used, with the possible exception of active duty examinees where USAFSAM aircrew norms may be applied. Clarification of profiles showing response biases, questionable validity, and/or unusual item endorsements will be conducted in individual test debriefing.

(c) Shipping Instructions

Forward all test materials as scored with annotations, interpretations, and impressions to the examining physician in your facility or MAIL DIRECTED TO

USAFSAM/EK BROOKS AFB TX 78235

and provide copy of letter of transmittal to the examining physician.

(d) <u>Psychometrics: Special Instructions</u>

(1) For the Cornell Index and MMPI, each subject is instructed:
 (a) to answer carefully <u>every</u> item; and (b)

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that wherever applicable, his responses should reflect personal experiences, beliefs, preferences, etc., <u>only</u> for the time period between his combat tour in SEA and the date of testing. These instruments are <u>not</u> to be group administered and a reasonable amount of privacy should be provided. These instruments should not be completed at the subject's overnight quarters nor anywhere else outside the supervised confines of the evaluative facility.

(2) If a subject's measured word recognition falls below the $\overline{6.5}$ Grade Level (Raw Score=40, Level II) according to the WRAT Reading subtest, the Cornell Index and MMPI are read aloud or administered via tape recording. In such cases, the subject retains the right to mark his answer sheet outside the view of the examiner or of others within hearing distance.

 $(\underline{3})$ All eleven subtests of the WAIS are administered, i.e., pro-rating of subtests is <u>not</u> allowed. The scoring of WAIS subtest items, and the operations of summing, transferring, and finding Raw Scores, Scaled Scores, and Tabled IQ values are double-checked for accuracy by the Psychologist in charge (or his/her appointed representative) before the raw data are forwarded to Brooks AFB.

 $(\underline{4})$ Precautions similar to those in $\Delta 3$ above are exercised in the scoring and other clerical tasks associated with the Halstead-Reitan, WMS I, WRAT, Cornell, and MMPI.

(5) For the Halstead-Reitan, use as the preferred, or dominant, hand the one which the subject uses most in writing. If in doubt, administer a "Name Writing Test", where the subject is simply asked to write his name in a normal manner as though signing a personal check. The examiner measures the time for each hand to perform, (without alerting <u>S</u> to the timing), and assigns dominance to the quickest hand.

 $(\underline{6})$ For the grip strength measure, report the average, in kilograms, of 3 brief, but maximum, squeezes of the dynamometer for the preferred and the non-preferred hands. Alternate hands between trials.

 $(\underline{7})$ The Psychologist in charge will conduct a one-to-one test debriefing with each subject to estimate the testby-test and overall accuracy and validity of the test results. A prepared form is provided for this purpose, and should be filled out completely before forwarding, with the subject's raw data, to Brooks AFB. If applicable, input from the testing technician utilized is encouraged. (3) Electrocardiogram

(a) A standard 12-lead scalar electrogram is required. If an arrhythmia is observed, a one minute rhythm strip is requested, in addition.

(b) <u>Mounting</u>: Mount the tracing in the usual manner of the laboratory for the recorder used.

(c) <u>Disposition</u>: Forward the mounted tracing and rhythm strip, if obtained, to the examining physician.

(d) Interpretation:

(1) The electrocardiograms will be interpreted by physicians in the USAF Central ECG Library and compared to previous individual ECG records in the case of rated (pilot or navigator) subjects.

(2) The interpretation and standard Central Library codes will be recorded on SAM Form 222 and forwarded to USAFSAM/ES.

(e) Disposition (USAF Central ECG Library):

(1) <u>Pilots and Navigators</u> - The original tracings will be microfisched and added to the individual's permanent record.

(2) <u>Enlisted Subjects</u> - The original tracings will be microfisched and a permanent record established for each individual.

- (4) Radiographic Examination
 - (a) Examination

A standard 14x17 in., standing, teleroentgenogram in the PA position using small nipple markers.

(b) Disposition

Forward the original film to the examining

physician or mail to

USAFSAM/EK Brooks AFB TX 78235

(c) Interpretation

USAFSAM/NGFR will interpret the teleroentgenogram and record the results on SAM Form 23. USAFSAM/NGAR will code the Radiologist's diagnosis (ICDA-9) and forward Sam Form 23 to USAFSAM/ES/

(5) Laboratory Procedures

(a) General Instructions; First Day

(1) The patient should report in the morning in a fasting state having had water only after midnight. The patient will have been requested to eat approximately 150 gms of carbohydrate each of the three preceding days and to consume no alcoholic beverages. Non-compliance is not a contraindication to drawing the blood specimens. However, a notation of extent of noncompliance should be made by the examining physician to aid in the interpretation of the results.

(2) The following is needed:

 (\underline{a}) Blood will be drawn only in the morning into a tube set-up consisting of the following: 4 large 15 ml red top clot tubes and 1 10-ml lavendar top EDTA tube.

(b) Label tubes with patient's full name, Social Security Number, date, and time of drawing.

(<u>c</u>) Perform routine hematology and sedimentation rate on the EDTA tube.

(d) Allow clot tubes to fully clot for at least 30 minutes. Centrifuge and separate hemolysis-free serum into screw-cap polypropylene tubes labeled with the patient's full name, Social Security Number, date, and time of drawing. Also label these tubes with the roman numeral I. Freeze tubes at -20° C as soon as possible (not to exceed 2 hours after drawing).

(e) After the drawing of the fasting specimens, administer 40 gms of glucose per square meter of body surface to the patient. Exactly 2 hours later draw one 7 ml red top clot tube. Alow tube to clot for 30 minutes, centrifuge, and separate hemolysis-free serum into a screw cap polypropylene tube. Label this tube with patient's full name, Social Security Number, date, and time of drawing. Label this tube "Ip.p." Freeze at -20° C as soon as possible not to exceed 2 hours after drawing.

 (\underline{f}) Ship all specimens frozen, packed in dry ice, by Federal Express-Priority one. Submit a patient list containing patient's full name, Social Security Number, and date of drawing. Address to:

> USAFSAM/NGP BLDG 125, Rm W-21 Brooks AFB, TX 78235

WARNING: DO NOT SHIP ON WEEKENDS, THURSDAY OR FRIDAY, OR ON ANY DAY PRIOR TO A FEDERAL HOLIDAY.

(3) The RANCH HAND II Protocol calls for a standard complete blood count, RBC indices, erythrocyte sedimentation rate, and routine urinalysis including a "dip stick" test for porphobilinogen and semen analysis. Since these tests must be done promptly, it is requested that the laboratory of the examining facility draw specimens and accomplish these procedures according to the laboratory's usual routine and forward the results to the examining physician at that facility.

(4) The RANCH HAND II Protocol calls for determination of delta-aminolevulinic acid and products of porphyrin metabolism. For these studies freeze $(-20^{\circ}C)$ a 100 ml aliquot of urine. The 100 cc urine aliquot must be acidified with 1 ml of glacial acetic acid. Collection of urine should be mid-morning of second day after blood for hormone analysis is drawn. Specific instructions for shipping these specimens will be supplied by USAFSAM/NGP.

(b) General Instructions; Second Day

Serum hormone levels should be determined from specimens collected on the morning of the second day. Hormonal levels appear to oscillate rapidly in a random fashion. Distributions drift with time suggesting diurnal variations and some are affected by nonfasting state. Therefore, the following instructions are critical:

 $(\underline{1})$ Patients should be fasting prior to drawing blood for hormone analysis.

 $(\underline{2})$ Exact time of each drawing should be recorded on each tube.

(3) One small clot tube $(7 \text{ ml}-without anticoagulant})$ should be drawn <u>every</u> 20 minutes for one hour. Patients should be kept at rest during the one-hour period. They should not smoke or drink stimulants (coffee or tea).

(4) RBCs should be separated from the serum within 2 hours of drawing the sample and the serum and the serum frozen as soon as possible at -20°C in 3 screw-top vials.

(5) Label each of 3 screw-top vials with time, date, and patients name followed by Roman numeral II.

(<u>6</u>) Ship specimens in dry ice in special containers by Federal Express-Priority One. Do not ship on Thursday, Friday, or the day before federal holidays.

- (c) Specific Tests
 - (1) Performed at the Examining Facility
 - (a) Hematocrit
 - (b) Hemoglobin
 - (c) RBC Indices
 - (d) While Blood Cell Count
 - (e) Platelet Count
 - (f) Erythrocyte Sedimentation Rate
 - (g) Urinalysis

<u>-(h-)</u>

-Semen Analysis (Number, Motility, Morphology)

- (2) Performed by USAFSAM Clinical Pathology Laboratory
 - (a) Blood Urea Nitrogen
 - (b) Fasting Plasma Glucose
 - (c) Creatinine
 - (<u>d</u>) 2-hour Post Prandial Plasma Glucose
 - (e) Differential Cortisol (0730 and 0930 hours)
 - (f) Cholesterol & HDL cholesterol
 - (g) Triglycerides
 - (h) SGOT
 - (i) SGPT
 - (j) GGTP
 - (k) Alkaline Phosphatase
 - (1) LDH
 - (m) Serum Protein Electrophoresis

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- (n) CPK
- (o) VDRL
- (3) Performed by USAFSAM Epidemiology Division Reference Laborator
 - (<u>a</u>) LH
 - (b) FSH
 - (c) Testosterone
 - (d) Thyroid Profile (RIA)
 - (<u>e</u>) Delta-aminolevulinic Acid
 - (f) Urine Porphyrins
- (<u>4</u>) Performed at USAFSAM if liver function studies are abnormal
 - (a) Anti-nuclear Antibody
 - (b) Hepatitis Antigens/Antibodies (A and B)
- (5) Performed if medical history indicates an increase in infectious diseases:
 - (a) Immuno electrophoresis
 - (b) Monilia Skin Test
 - (<u>c</u>) Quantitative Immunoglobulin Determinations
 - (d) Rationale for Laboratory Procedures

(1) Studies on the toxicity of TCDD in animals have shown that the following organ systems are damaged:

(<u>a</u>) Liver: Hepatic necrosis, liver enzyme changes, hypoproteinemia, hypercholesterolemia, hypertriglyceridemia.

(b) Reticuloendothelial System: Thymic atrophy, altered cellular immunity, decreased lymphocyte counts.

(<u>c</u>) Hemopoietic System: Anemia, thrombocytopenia, leukopenia, pancytopenia.

Hemorrhage and (d) Endocrine System: atrophy of adrenal cortex, hypothyroidism. (e) Renal: Increase in blood urea nitrogen. In addition, statistically significant increases in hepatocellular carcinomas (liver) and squamocellular carcinomas of the lung were found. (2) Studies on the toxic effects of TCDD in man have shown that the following organ systems are damaged: (a) Skin: Chloracne, hirsutism. (b) Liver: Porphyria cutanea tarda. Increased levels of transaminase and of GGTP. Enlarged, tender liver, hyperlipidemia. (c) Renal: Hemorrhagic cystitis, focal Pyelonephritis. (d) Neuromuscular System: Asthenia. apathy, fatigue, anorexia, weitht loss, sleep i.e., headache, disturbances, decreased learning ability, decreased memory. dyspepsia, sweating, muscle pain, joint pain and sexual dysfunction. (e) Endocrine System: Hypothyroidism. (3) Based upon the reports of toxic effects in animal and human exposures, the following organ panels are recommended: (a) Hemopoietic (b) Reticuloendothelia] (c) Rena 1 (d) Endocrine (e) Neuromuscular (4) Hemopoietic screening should include: (a) Hematocrit (<u>b</u>) Hemoglobin (c) **RBC** indices

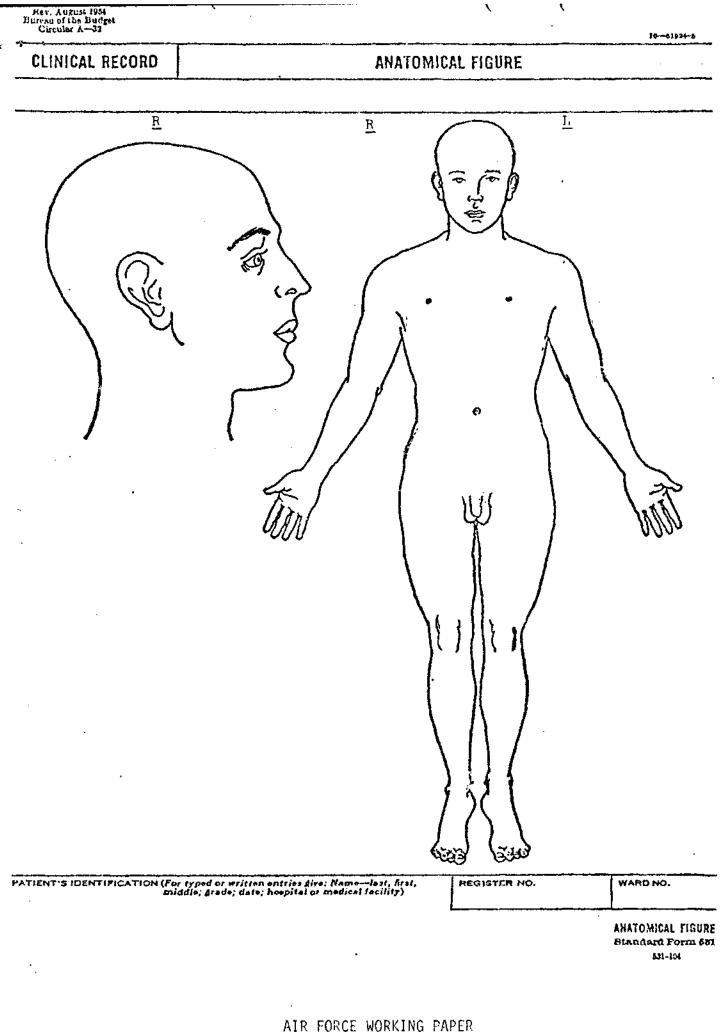
- (d) Erythrocyte sedimentation rate
- (e) Platelet count
- (5) Reticuloendothelial system:
 - (a) White blood cell count
 - (b) Differential
 - (c) Serum protein electrophoresis

 (\underline{d}) Selective use of skin testing, immunoelectrophoresis and quantative immunoglobulin determination

- (6) Hepatic screen:
 - (a) SGOT
 - (b) SGPT
 - (c) GGTP
 - (d) Alkaline phosphatase
 - (e) LDH
 - (f) Cholesterol
 - (g) HDL cholesterol
 - (h) Triglyceride
 - (i) Urine prophyrins
 - (j) Urine porphobilinogen
- (7) Renal screen:
 - (<u>a</u>) Urinalysis
 - (b) BUN
 - (<u>c</u>) Creatinine
- (8) Endocrine screen
 - (<u>a</u>) Differential cortisol (0730 and 0930 hours)

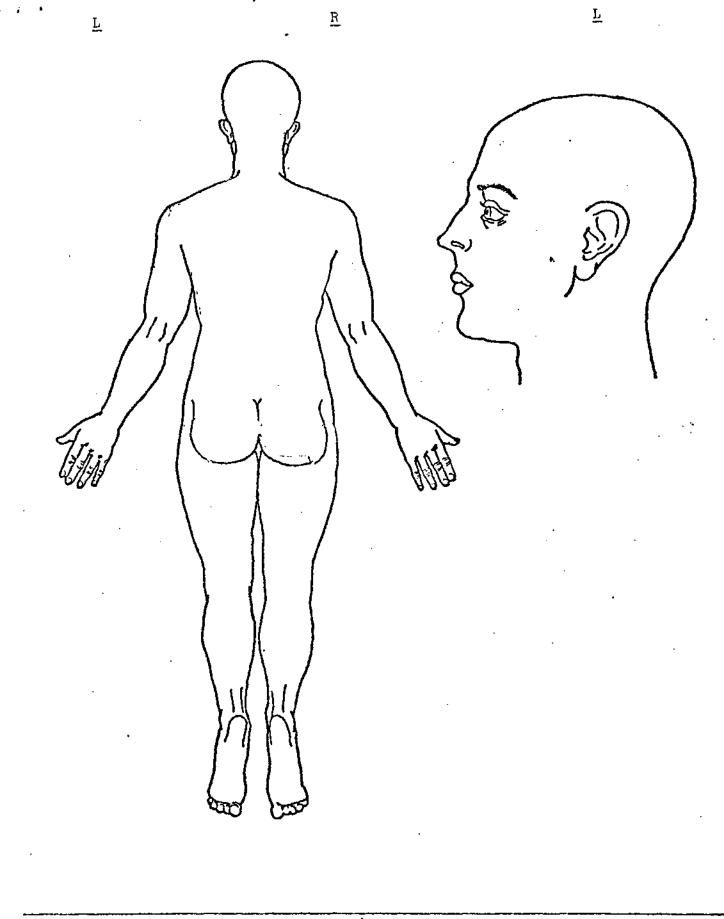
E. Forms

Anatomical Figure (Anterior) Anatomical Figure (Posterior) Nerve Conduction Velocities Psychometric De-Briefing Form Radiographic Data Electrocardiographic Studies



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1. Ulnar (one side	$\leq only) / R$	L Elbow Above Below
· ·		Normal Values for Laboratory
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		Normal Values for Laboratory
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Stm. Curr. /	<u>/ / /</u> mA	<u>/_/_/</u> /

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Ranch Hand II: Psychometric De-Briefing Form

Subject:				<u> </u>
	Name	Test Date	Eval Faci	lity Handedness
Psychologist/ De-Briefer :				Yes No.
	Name	Title D	egree Clin,	/Couns Cert/Lic
Testing <u>Technician</u> :	· · · · · · · · · · · · · · · · · · ·			
	Same as above	Name	Degree	Test/Experience (Yrs)

Instructions

In the appropriate column below, indicate the test-by-test validity of the psychometric results based upon the Examiner's observations of the subject during testing and upon the Psychologist's evaluation of the data in test de-briefing with the subject. Use the numbered factors below to indicate the reason(s) for questionable validity among any of the data. For datum thought to be of questionable validity, also provide an estimate of the subject's "true" score or result. Forward the completed form with the subject's raw data.

Reasons for Questionable Validity

- 1. Poor reading comprehension
- 2. Fatigue
- Neg attitude, angry, marginal cooperator
- 4. Careless, hurried responses
- 5. Examiner Error

- 6. Exaggeration of complaints ("fake bad")
- 7. Minimizing complaints ("fake good")
- 8. Disorganized personality (Psychotic)
- 9. Physically ill (flu, venipuncture effects, etc)
- 10. Other (Specify_____)

	<u>Test Score</u>	<u>Valid Results</u>	Reason(s) for Questionably <u>Valid Results</u>	Est of "True" Score/Result
1.	WAIS VIQ PIQ FSIQ			
2. 3.	WRAT Reading Halstead-Reitan Category Test			

Ranch Hand II: Psychometric De-Briefing Form Continued

	<u>Test Score</u>	Valid <u>Results</u>	Reason(s) for Questionably Valid Results	Est of "True" <u>Score/Result</u>
4.	Tactual Performance Test Preferred Hand Non-Preferred Hand Both Hands Memory Localization Speech-Sounds Perception Seashore Rhythm Finger Tapping Preferred Hand Non-Preferred Hand Trail Making Test Part A Part B Grip Strengths Preferred Hand Non-Preferred Hand WMS I Logical Mem (immed)			
5. 6.	Visual Repro (immed) Associate Lrng Logical Mem (delayed) Visual Repro (delayed) Cornell Index MMPI (overall rating of protocol)			WNL or ONL

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PROJECT RANCH HAND II

EXAMINER'S HANDBOOK

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A. General Instructions

Project RANCH HAND II is a multiyear effort to determine whether or not C-123 aircrew members who were engaged in the aerial spraying of herbicides in Vietnam have developed significant adverse health effects from that exposure. Detailed surveys of the world's literature have been used in designing the history questionnaires, physical examination protocol, and laboratory procedures.

This phase of Project RANCH HAND II involves a cross sectional study of the subject's health at the time of examination. It is important that examiners remain unaware of the subject's status as a RANCH HAND participant or as a control subject. The physician examiner is tasked to examine and objectively record his findings. The examining physician is not, and cannot be expected to arrive at any definitive diagnosis as the full history and laboratory results are not available to him. The compilation and analysis of data will be performed by the study investigators at Brooks Air Force Base, Texas. They will notify the subject and the physician of his choice of the results of the examination.

The physicians performing examinations for Project RANCH HAND II should be aware that the report of examination will become a permanent record. This report will be referred to not only in the near future as the cross sectional study is analyzed, but at the time of the next review of the subject in the follow-up phases of Project RANCH HAND. These examinations will define the health status of the subjects at a point in time, and to establish the presence of physical findings, if any exist. After statistical review of the study groups, these findings may permit definition of a chronic effect due to exposure. An inaccurate examination may lead to falacious study results in two ways: a presumed syndrome may be defined which does not in fact exist, or a syndrome which in fact exists may not be defined with enough validity to warrant further actions.

The examining physician is responsible for recording a complete and detailed report of the physical examination. In this role, the examining physician is tasked with collecting evidence of the presence or absence of physical signs of abnormality only. Formulation of impressions is not requested nor desired. All items on the physical examination report form must be completed. It is imperative that the physician make such additional remarks as may be required to adequately describe existing physical and mental impairments. The examining physician must avoid an expression of opinion regarding the interpretation of any findings particularly with regards to possible etiologies. If, during the examination, the physician discovers evidence of acute serious illness requiring immediate treatment, the normal emergency or urgent care procedures of the medical facility would apply. If during the examination, the examining physician finds evidence of present illness requiring further medical attention, he should so state to the subject and

offer to forward or have forwarded pertinent information to the subject's physician. A clear record of any such advice and treatment should be recorded. The ultimate value of the RANCH HAND II Study will lie in complete, accurate and, whenever possible, quantitative data permitting the most stringent and powerful statistical analysis. For that reason, the physical examination protocol requires exact measurements in many instances, and the use of defined meanings of semiquantitative indicators in other places.

B. Conduct of the Examination

(1) On arrival at the examining facility the subject should be briefed on the appointments which have been arranged, their times, and locations.

(2) Collation and forwarding of examination results

A checklist for the mailing of data will be provided. It should be retained by the office primarily responsible (OPR) for the Project RANCH HAND II examination and used to ascertain that all necessary items have been completed and received, or have been directly forwarded by the section performing the examinations. When the OPR for the examinations is ready to forward all materials, the checklist for mailing should be endorsed with the date of mailing as a letter of transmittal and included in the package of material to be mailed to USAFSAM/EK. Brooks AFB TX 78235.

(3) Forms for individual examinations and procedures

The blank forms included for various examinations and procedures may be carried by the patient so as to be available to the examiner or to the laboratory, or to the department of radiology, as the patient reports for his examinations in those functions. The forms pertaining to the specific function may be withdrawn from the patient's examination package and later returned to the office of primary responsibility.

SECTION		PHYSICAL EXAMINATION		· · · · · · · · · · · · · · · · · · ·	SUBJECT NUMBER
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17. NERVOUS SYSTEM - SEE	ATTACHE	D FORMS	• •			-
18. HEART AND OTHER OBSE	RVATIONS			· · · · · · · · · · · · · · · · · · ·		
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CLINICAL RECORD NEUROLOGICAL EXAMINATION HEAD AND NECK - Normal to Palpations/Inspection ____Y ___N Specify Scar ____ Asymmetry **[7** Depression **[7**] Carotid Bruit 7No 7R 7L Neck Range of Motion / Normal or Decreased to / Left / Right / Forward / Backward TRUNK MOTOR SYSTEM - Handedness Right /7 Left /7 Gait [] Normal or [] Broad Based [] Ataxic []Small Stepped /]Other-Specify Associated Movements / Arm Swing / Normal or Abnormal / 7R / 7L Muscle Status (strength, tone, volume, tenderness, fibrillations) Bulk [7 Normal /7 Abnormal Tone Upper Extremities []Normal or []Increased []Decreased /7Right /7Left Lower Extremities [7]Normal or [7]Increased [7]Decreased /7Right /7Left Strength - Distal wrist extensors / 7Normal / 7Decreased Ankle/Toe Dors/Flexors / Normal / Decreased / 7R / 7L Proximal Deltoids //Normal //Decreased //R //L Hip Flexors //Normal //Decreased //R //L Abnormal Movements (tremors, tics, choreas, etc.) Fasiculations /7No /7Yes (1-4+) Tenderness TNo TYes (1-4+) Tremor /7No /7Yes - Specify Upper Extremity [7R [7] [7] Resting [7] Essential [7] Intention Lower Extremity / R / L Other Coordination (a) Equilibratory - Eyes Open Eyes Closed - Romberg ____Positive (Abnormal) ____Negative (Normal) **Right Foot** Left Foot (b) Nonequilibratory (F to N; F to F; H to K) Finger-to-nose-to-finger //Normal //Abnormal //Right //Left //Both
Heel-Knee-Shin //Normal //Abnormal //Right //Left //Both
Succession Movements (including check, rebound, posture-holding) (c) If indicated, check //Normal //Abnormal //R //R Rapidly alternative movements / Normal / Abnormal / R / L / Both Skilled Acts (a) Praxis (b) Handwriting. If indicated, / Normal / Abnormal Speech (articulation, aphasia, agnosia) Grossly //Normal (c) [7] Abnormal - Specify Dysarthria [7] Aphasia 🔼 AIR FORCE WORKING PAPER

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Ptosis R/7 L /7 V Motor R Clench Jaw - Symmetric / Deviated / R/ L/ R Normal \square Abnormal \square $V_1 \square$ $V_2 \square$ $V_3 \square$ L Normal \square Abnormal \square $V_1 \square$ $V_2 \square$ $V_3 \square$ Sensory **Corneal Reflex** R L VII Motor R Normal smile / Yes / No Palpebral Fissure / Yes / No L Normal smile ___Yes __No Palpebral Fissure ___Yes __No IX Palate and Uvula X Movement Normal [7] Deviation to [7R [7]L Palatal Reflex R / Normal / Abnormal L [7Norma] [7Abnorma] XII Tongue-Protruded-Central [7 R [7 L [7 Atrophy 7No 7Yes MENTAL STATUS (alert, clear, cooperative, etc.) Gross abnormalities: ____No /7Yes - Specify

SUMMARY OF POSITIVE FINDINGS Objective

Subjective

Diagnostic Impression

Date

Signature

D. Special Procedures

(1) Nerve Conduction Velocities

(a) These studies have been determined to be an important parameter in long-term follow-up studies of persons thought to have been exposed to Herbicide Orange Components.

(b) The Nerve Conduction Velocities should be performed by a physician or by a specialty qualified technician under the supervision of a physician trained in neurophysiological methods.

- (c) <u>Specific NCVs</u> (See form included in F. Below)
 - (1) Ulnar Nerve (one side only)
 - (a) motor (above elbow, below elbow)
 - (b) values recorded
 - (i) distal latency
 - (11) NCV
 - (2) <u>Peroneal Nerve</u> (one side only)

(a) motor

- (b) values recorded
 - (i) distal latency
 - (ii) NCV
- (3) <u>Sural Nerve</u> (one side only)
 - (<u>a</u>) sensory: orthodromic
 - (b) values recorded: NCV

(d) <u>Methods</u>

PERONEAL NERVE

 $(\underline{1})$ Active electrode is placed over the extensor digitorum brevis and reference over the little toe. Stimulating electrodes are placed over anterior distal leg 8 cm proximal to active electrode. Proximal site is distal to head of fibula. If entrapment is suspected at fibular head use a stimulation site of 12-18 cm more proximal to the fibular head.

Anomalous innervation to the extensor digitorum brevis occurs in 1/5 patients (at least partially). Identified by inability to evoke a muscle action potential when stimulating at anterior ankle or a different shape (smaller) potential when stimulating here. This accessory nerve causes posterior to lateral malleolus so cathode should be placed here.

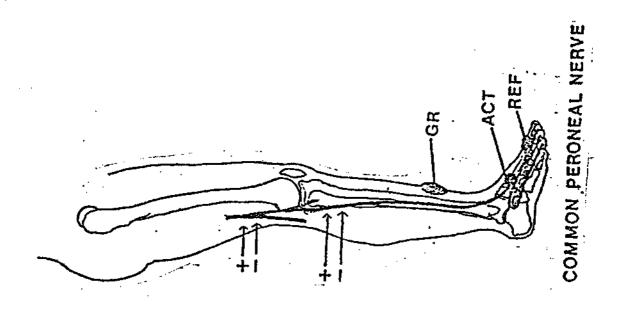
NORMAL VALUES

49.9 \pm 5.9 M/sec Distal latency: 4.5 \pm .8 ms

Proximal latencies have been determined for use in below the knee amputees, and neuromuscular diseases where extensor digitorum brevis action potential cannot be elicited. Active electrode is placed 1/2 way down leg over middle of dorsiflexor muscle group and stimulation at fibular head.

NORMAL VALUES

5.5 - 7.2 ms (N = 217)

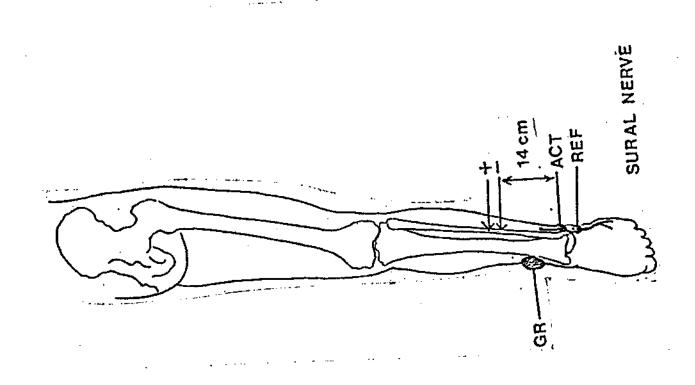


SURAL NERVE

 $(\underline{2})$ Active and recording electrodes are placed under lateral malleolus on lateral aspect of ankle. Sural nerve is stimulated as it pierces the gastrocnemius fascia just lateral to the midline of posterior distal calf, 10-18 cm proximal to active electrode. If leg is cold - a clue is prolonged latency of peroneal nerve - determine temperature. Subtract .1 ms (latency of activation) from the observed latency and divide into the distance.

NORMAL VALUES (after LaFratta)

Age	(To Peak)
20-29	44 ± 2.5 M/sec
30-39	38.80 ± 3.3 M/sec
40-49	36.70 ± 3.7 M/sec
50-59	37.20 ± 3.0 M/sec
60 a over	35.00 ± 3.8 M/sec



ULNAR NERVE

MOTOR CONDUCTION

(3) Active electrode is placed over center of abductor digiti quinti; reference over proximal phalanx fifth digit. Stimulation (cathode) just radial to tendon of flexor carpi ulnaris 8 cm proximal to active electrode. Proximal site of stimulation should be just below ulnar groove and 18 cm proximal to ulnar groove on medial aspect of humerus.

<u>N.B.</u>: Elbow should be flexed to 70 degrees during procedure of stimulation and measurement to make more precise the actual length of ulnar nerve. More proximal stimulation sites include supraclavicular and C-8 root (see median nerve).

SENSORY CONDUCTION

Antidromic - ring electrodes over fifth digit separated by 4 cm. N.B. motor artifact may be interfering. Stimulate 14 cm proximal to active electrode at same site as motor stimulation.

Orthodromic - reverse stimulation and recording electrodes. More proximal sites of stimulation may also be done.

NORMAL VALUES

57 \pm 4.7 M/sec - motor forearm segment 62.7 \pm 5.5 M/sec - motor across elbow segment 56.7 \pm 4.2 M/sec - sensory orthodromic (to peak) 54.9 \pm 3.9 M/sec - sensory antidromic (to peak)

Distal Latency:

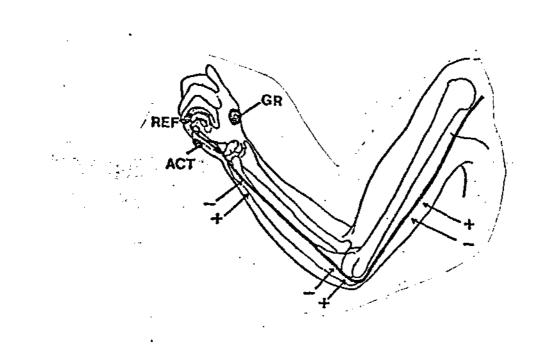
Motor: $3.7 \pm .3$ Sensory: $3.0 \pm .25$ Antidromic (peak) $3.0 \pm .25$ Orthodromic (peak)

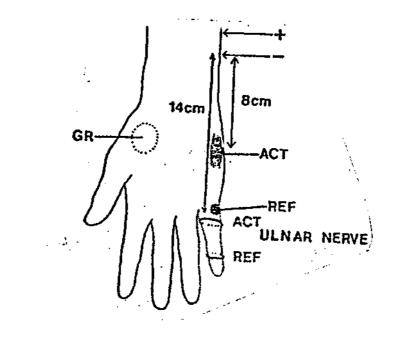
Muscle AP 8-20 mV

Sensory AP 15-50 mV

ADDENDUM

For deep branch surface recording electrode should be over adductor pollicus (i.e. just medial to thenar eminence on palmar surface of web space). Additional latency is .5 ms.





(e) Disposition

Forward the recorded results on the form attached to the examination package to the examining physician.

(2) Psychological Battery

(a) <u>General</u>

(1) This battery yields objective numerical data, and is well-standardized and clinically validated. The individual tests were chosen to insure an adequate analysis of one of the major alleged manifestations of herbicide toxicity. Each test either validates one of the other tests, or is considered to be a "definitive" test for analyis of a suspected psycho/neuropathic effect.

(2) Compared to the general civilian population, characteristic response tendencies are observed on the MMPI and Cornell Index among active duty aircrewmen being evaluated in an aeromedical setting. It is also important to consider the effect that pending retirement has exerted on the reporting of medical history and symptomatology. This may also alter responses to psychological testing.

(3) The battery requires approximately 5-1/2 to 6-3/4 hours to administer, depending on the speed of the examinee. An additional 1 to 2 hours of scoring and other clerical tasks will be required. Since test debriefing to clarify unusual performances, response biases, etc., is a crucial part of the psychologic evaluation, it is recommended that testing begin and be completed as early as possible during each examinee's stay at his respective evaluative facility.

(b) <u>Specific Tests</u>

(<u>1</u>) <u>Wechsler Adult Intelligence Scale</u> (<u>WAIS</u>): 60-75 minute individually-administered collection of verbal and nonverbal intellectual measures; also useful for clinical inferences when combined with the neuropsychological battery below.

(2) <u>Reading subtest of the Wide Range</u> <u>Achievement Test (WRAT)</u>: 10-minute individually-administered measure of word recognition ability. Important so as to rule-out reading inefficiency should response to personality instruments below be of questionable validity (e.g., high F Scale on MMPI).

(3) <u>Halstead-Reitan Neuropsychological Test</u> <u>Battery</u>: 150-180 minute individually-administered collection of brain behavior relationship measures for establishing the functional integrity of the cerebral hemispheres. The battery must include the following subtests: Category, Tactual performance, Speech-Sounds,

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Seashore Rhythm, Finger Tapping, Trail Making, and Grip Strengths. The Aphasia Screening and Sensory-Perceptual Exams are considered optional in view of their redundancy with the clinical neurologic exam included in this project. Individualized test debriefing is conducted to clarify test performances in the WAIS and Neuropsychological Battery.

(4) <u>Three subtests of the Wechsler Memory</u> <u>Scale I (WMS I)</u>: 30-minute individually-administered measures of immediate and delayed recall of verbal and visual materials. The Logical Memory, Associate Learning and Visual Reproduction subtests are to be administered in the standard, immediate-recall fashion initially. After 30 minutes has elapsed, the examinee is asked, without prior alerting, to recall as much as he can about the Logical Memory and Visual Reproduction subtest stimuli. Standard scoring is used for both test-retest administrations.

(5) <u>Cornell Index (CI)</u>: 10-15 minute selfadministered and standardized neuropsychiatric symptom and complain inventory, including items involving asthenia, depression, anxiety, fatigue, and GI symptoms in lay language. Endorsement of items are to be explored and clarified in test-debriefing.

(6) <u>Minnesota Multiphasic Personality</u> <u>Inventory (MMPI)</u>: 60 to 90 minute self administered clinical psychiatric screening instrument; also capable of estimating response biases (e.g., "fake good," or "fake bad"). The shortened version of Form R (i.e., items 1 to 399) may be substituted for the 566-item Long Form. Standard scoring and Minnesota norms are to be used, with the possible exception of active duty examinees where USAFSAM aircrew norms may be applied. Clarification of profiles showing response biases, questionable validity, and/or unusual item endorsements will be conducted in individual test debriefing.

(c) Shipping Instructions

Forward all test materials as scored with annotations, interpretations, and impressions to the examining physician in your facility or MAIL DIRECTED TO

USAFSAM/EK BROOKS AFB TX 78235

and provide copy of letter of transmittal to the examining physician.

(d) Psychometrics: Special Instructions

(1) For the Cornell Index and MMPI, each subject is instructed:
 (a) to answer carefully <u>every</u> item; and (b)

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that wherever applicable, his responses should reflect personal experiences, beliefs, preferences, etc., <u>only</u> for the time period between his combat tour in SEA and the date of testing. These instruments are <u>not</u> to be group administered and a reasonable amount of privacy should be provided. These instruments should not be completed at the subject's overnight quarters nor anywhere else outside the supervised confines of the evaluative facility.

(2) If a subject's measured word recognition fails below the 6.5 Grade Level (Raw Score=40, Level II) according to the WRAT Reading subtest, the Cornell Index and MMPI are read aloud or administered via tape recording. In such cases, the subject retains the right to mark his answer sheet outside the view of the examiner or of others within hearing distance.

(3) All eleven subtests of the WAIS are administered, i.e., pro-rating of subtests is <u>not</u> allowed. The scoring of WAIS subtest items, and the operations of summing, transferring, and finding Raw Scores, Scaled Scores, and Tabled IQ values are double-checked for accuracy by the Psychologist in charge (or his/her appointed representative) before the raw data are forwarded to Brooks AFB.

 $(\underline{4})$ Precautions similar to those in $\Delta 3$ above are exercised in the scoring and other clerical tasks associated with the Halstead-Reitan, WMS I, WRAT, Cornell, and MMPI.

(5) For the Halstead-Reitan, use as the preferred, or dominant, hand the one which the subject uses most in writing. If in doubt, administer a "Name Writing Test", where the subject is simply asked to write his name in a normal manner as though signing a personal check. The examiner measures the time for each hand to perform, (without alerting <u>S</u> to the timing), and assigns dominance to the quickest hand.

 $(\underline{6})$ For the grip strength measure, report the average, in kilograms, of 3 brief, but maximum, squeezes of the dynamometer for the preferred and the non-preferred hands. Alternate hands between trials.

(7) The Psychologist in charge will conduct a one-to-one test debriefing with each subject to estimate the testby-test and overall accuracy and validity of the test results. A prepared form is provided for this purpose, and should be filled out completely before forwarding, with the subject's raw data, to Brooks AFB. If applicable, input from the testing technician utilized is encouraged.

(3) Electrocardiogram

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(a) A standard 12-lead scalar electrogram is required. If an arrhythmia is observed, a one minute rhythm strip is requested, in addition.

(b) <u>Mounting</u>: Mount the tracing in the usual manner of the laboratory for the recorder used.

(c) <u>Disposition</u>: Forward the mounted tracing and rhythm strip, if obtained, to the examining physician.

(d) Interpretation:

(1) The electrocardiograms will be interpreted by physicians in The USAF Central ECG Library and compared to previous individual ECG records in the case of rated (pilot or navigator) subjects.

(2) The interpretation and standard Central Library codes will be recorded on SAM Form 222 and forwarded to USAFSAM/ES.

(e) Disposition (USAF Central ECG Library):

(1) <u>Pilots and Navigators</u> - The original tracings will be microfisched and added to the individual's permanent record.

(2) Enlisted Subjects - The original tracings will be microfisched and a permanent record established for each individual.

(4) Radiographic Examination

(a) Examination

A standard 14x17 in., standing, teleroentgenogram in the PA position using small nipple markers.

(b) Disposition

Forward the original film to the examining

physician or mail to

USAFSAM/EK Brooks AFB TX 78235

(c) Interpretation

USAFSAM/NGFR will interpret the teleroentgenogram and record the results on SAM Form 23. USAFSAM/NGAR will code the Radiologist's diagnosis (ICDA-9) and forward Sam Form 23 to USAFSAM/ES/

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(5) Laboratory Procedures

(a) General Instructions; First Day

(1) The patient should report in the morning in a fasting state having had water only after midnight. The patient will have been requested to eat approximately 150 gms of carbohydrate each of the three preceding days and to consume no alcoholic beverages. Non-compliance is not a contraindication to drawing the blood specimens. However, a notation of extent of noncompliance should be made by the examining physician to aid in the interpretation of the results.

(2) The following is needed:

 (\underline{a}) Blood will be drawn only in the morning into a tube set-up consisting of the following: 4 large 15 ml red top clot tubes and 1 10-ml lavendar top EDTA tube.

 (\underline{b}) Label tubes with patient's full name, Social Security Number, date, and time of drawing.

(c) Perform routine hematology and sedimentation rate on the EDTA tube.

(d) Allow clot tubes to fully clot for at least 30 minutes. Centrifuge and separate hemolysis-free serum into screw-cap polypropylene tubes labeled with the patient's full name, Social Security Number, date, and time of drawing. Also label these tubes with the roman numeral I. Freeze tubes at -20° C as soon as possible (not to exceed 2 hours after drawing).

(e) After the drawing of the fasting specimens, administer 40 gms of glucose per square meter of body surface to the patient. Exactly 2 hours later draw one 7 ml red top clot tube. Alow tube to clot for 30 minutes, centrifuge, and separate hemolysis-free serum into a screw cap polypropylene tube. Label this tube with patient's full name, Social Security Number, date, and time of drawing. Label this tube "Ip.p." Freeze at -20°C as soon as possible not to exceed 2 hours after drawing.

 (\underline{f}) Ship all specimens frozen, packed in dry ice, by Federal Express-Priority one. Submit a patient list containing patient's full name, Social Security Number, and date of drawing. Address to:

> USAFSAM/NGP BLDG 125, Rm W-21 Brooks AFB, TX 78235

WARNING: DO NOT SHIP ON WEEKENDS, THURSDAY OR FRIDAY, OR ON ANY DAY PRIOR TO A FEDERAL HOLIDAY.

(3) The RANCH HAND II Protocol calls for a standard complete blood count, RBC indices, erythrocyte sedimentation rate, and routine urinalysis including a "dip stick" test for porphobilinogen and semen analysis. Since these tests must be done promptly, it is requested that the laboratory of the examining facility draw specimens and accomplish these procedures according to the laboratory's usual routine and forward the results to the examining physician at that facility.

(4) The RANCH HAND II Protocol calls for determination of delta-aminolevulinic acid and products of porphyrin metabolism. For these studies freeze $(-20^{\circ}C)$ a 100 ml aliquot of urine. The 100 cc urine aliquot must be acidified with 1 ml of glacial acetic acid. Collection of urine should be mid-morning of second day after blood for hormone analysis is drawn. Specific instructions for shipping these specimens will be supplied by USAFSAM/NGP.

(b) General Instructions; Second Day

Serum hormone levels should be determined from specimens collected on the morning of the second day. Hormonal levels appear to oscillate rapidly in a random fashion. Distributions drift with time suggesting diurnal variations and some are affected by nonfasting state. Therefore, the following instructions are critical:

 $(\underline{1})$ Patients should be fasting prior to drawing blood for hormone analysis.

 $(\underline{2})$ Exact time of each drawing should be recorded on each tube.

(3) One small clot tube (7 ml--without anticoagulant) should be drawn <u>every</u> 20 minutes for one hour. Patients should be kept at rest during the one-hour period. They should not smoke or drink stimulants (coffee or tea).

 $(\underline{4})$ RBCs should be separated from the serum within 2 hours of drawing the sample and the serum and the serum frozen as soon as possible at -20°C in 3 screw-top vials.

(5) Label each of 3 screw-top vials with time, date, and patients name followed by Roman numeral II.

(6) Ship specimens in dry ice in special containers by Federal Express-Priority One. Do not ship on Thursday, Friday, or the day before federal holidays.

- (c) Specific Tests
 - (1) Performed at the Examining Facility
 - (a) Hematocrit
 - (b) Hemoglobin
 - (c) RBC Indices
 - (d) While Blood Cell Count
 - (e) Platelet Count
 - (f) Erythrocyte Sedimentation Rate
 - (g) Urinalysis

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- h) -Semen Analysis (Number, Motility, - Morphology)
- (2) Performed by USAFSAM Clinical Pathology Laboratory
 - (a) Blood Urea Nitrogen
 - (b) Fasting Plasma Glucose
 - (<u>c</u>) Creatinine
 - (<u>d</u>) 2-hour Post Prandial Plasma Glucose
 - (e) Differential Cortisol (0730 and 0930 hours)
 - (f) Cholesterol & HDL cholesterol
 - (g) Triglycerides
 - (h) SGOT
 - (1) SGPT
 - (j) GGTP
 - (k) Alkaline Phosphatase
 - (1) LDH
 - (m) Serum Protein Electrophoresis

- (n) CPK
- (o) VDRL
- (3) Performed by USAFSAM Epidemiology Division Reference Laborator
 - (a) LH
 - (b) FSH
 - (c) Testosterone
 - (d) Thyroid Profile (RIA)
 - √(e) Delta-aminolevulinic Acid
 - (f) Urine Porphyrins
- (<u>4</u>) Performed at USAFSAM if liver function studies are abnormal
 - (a) Anti-nuclear Antibody
 - (b) Hepatitis Antigens/Antibodies (A and B)
- (5) Performed if medical history indicates an increase in infectious diseases:
 - (a) Immuno electrophoresis
 - (b) Monilia Skin Test
 - (c) Quantitative Immunoglobulin Determinations
 - (d) Rationale for Laboratory Procedures

(1) Studies on the toxicity of TCDD in animals have shown that the following organ systems are damaged:

(a) Liver: Hepatic necrosis, liver enzyme changes, hypoproteinemia, hypercholesterolemia, hypertriglyceridemia.

(b) Reticuloendothelial System: Thymic atrophy, altered cellular immunity, decreased lymphocyte counts.

 (\underline{c}) Hemopoietic System: Anemia, thrombocytopenia, leukopenia, pancytopenia.

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(ď) Endocrine System: Hemorrhage and atrophy of adrenal cortex, hypothyroidism. (e) Renal: Increase in blood urea nitrogen. In addition, statistically significant increases in hepatocellular carcinomas (liver) and squamocellular carcinomas of the lung were found. Studies on the toxic effects of TCDD in (2) man have shown that the following organ systems are damaged: (a) Skin: Chloracne, hirsutism. (b) Liver: Porphyria cutanea tarda. Increased levels of transaminase and of GGTP. Enlarged, tender liver, hyperlipidemia. (c) Renal: Hemorrhagic cystitis, focal Pyelonephritis. (d) Neuromuscular System: Asthenia. i.e., headache, apathy, fatigue, anorexia, weitht loss, sleep decreased learning ability, decreased memory, disturbances, dyspepsia, sweating, muscle pain, joint pain and sexual dysfunction. (e) Endocrine System: Hypothyroidism. (3) Based upon the reports of toxic effects in animal and human exposures, the following organ panels are recommended: (a) Hemopoietic (b) **Reticuloendothelial** (c) Renal (d) Endocrine (e) Neuromuscular

- (4) Hemopoietic screening should include:
 - (<u>a</u>) Hematocrit
 - (b) Hemoglobin
 - (c) RBC indices

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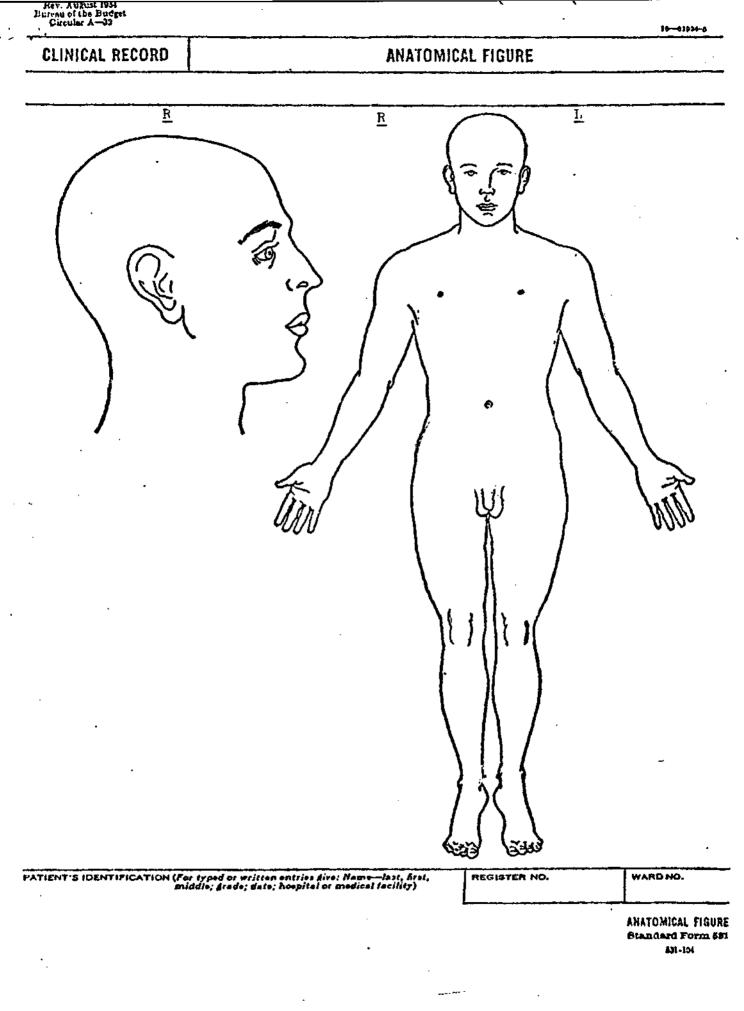
- (d) Erythrocyte sedimentation rate
- (e) Platelet count
- (5) Reticuloendothelial system:
 - (a) White blood cell count
 - (b) Differential
 - (c) Serum protein electrophoresis

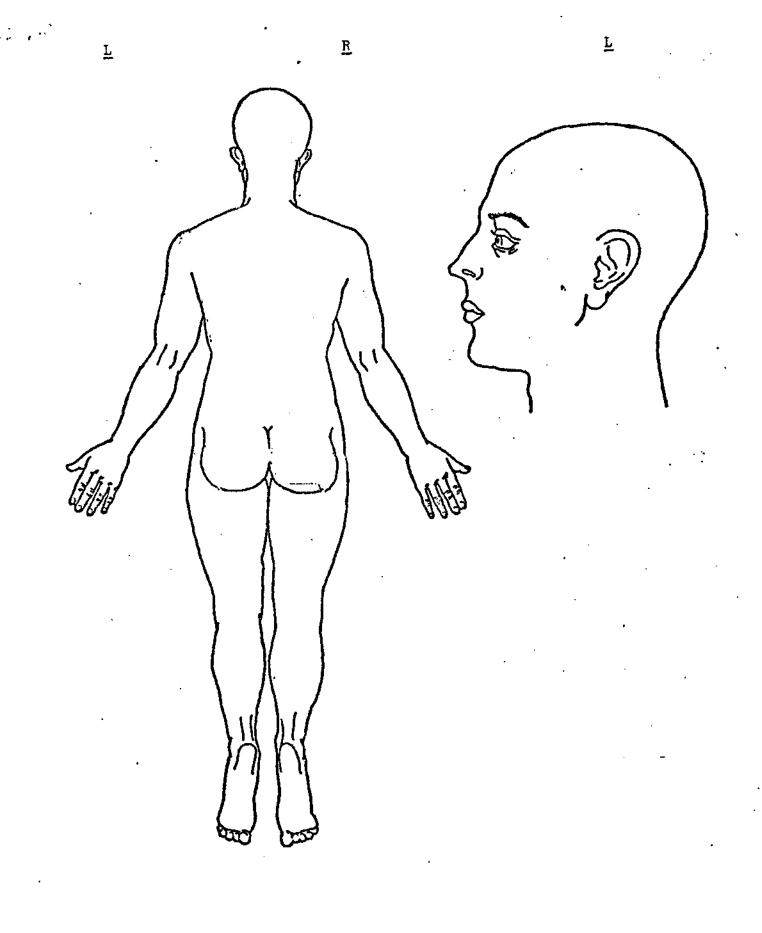
(d) Selective use of skin testing, immunoelectrophoresis and quantative immunoglobulin determination

- (6) Hepatic screen:
 - (a) SGOT
 - (b) SGPT
 - (c) GGTP
 - (d) Alkaline phosphatase
 - (e) LDH
 - (f) Cholesterol
 - (g) HDL cholesterol
 - (h) Triglyceride
 - (i) Urine prophyrins
 - (j) Urine porphobilinogen
- (7) Renal screen:
 - (a) Urinalysis
 - (b) BUN
 - (c) Creatinine
- (8) Endocrine screen
 - (a) Differential cortisol (0730 and 0930 hours)

E. Forms

Anatomical Figure (Anterior) Anatomical Figure (Posterior) Nerve Conduction Velocities Psychometric De-Briefing Form Radiographic Data Electrocardiographic Studies





U.S. GOVERNMENT PRINTING OFFICE (12-0-0-711-787

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1. Ulnar (one side only) / R	
	Normal Values for Laboratory
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Distance / / / / mm	<u>/ / / /</u>
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Stm. Curr. / / / mA	<u> </u>
2. Peroneal (one side only)R	<u> </u>
	Normal Values for Laboratory
Latency / / /./ / ms	<u>/ / /./ /</u>
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N.C.V. <u>/ / /./ /</u> m/s	<u>/ / /./ /</u>
Stm. Curr. <u>/ / /</u> mA	<u> </u>
3. Sural (one side only)R	7L (If unobtainable, Median or Ulnar Sensory recommend)
	Normal Values for Laboratory
Latency / / /./ / ms	<u> . </u>
Distance / / / mm	
N.C.V. / / /./ / m/s	<u>/ / /./ /</u>
Stm. Curr. / / / mA	<u> </u>

Ranch Hand II: Psychometric De-Briefing Form

Subject:			·····	<u> </u>
-	Name	Test Da	ite Eval Fac	ility Handedness
Psychologist/ De-Briefer _:				Yes No n/Couns Cert/Lic
· ·	Name	Title	Degree Cli	n/Couns Cert/Lic
Testing Technician:	· · · · · · · · ·			
- <u></u>	Same as above	Name	Degree	Test/Experience (Yrs)

Instructions

In the appropriate column below, indicate the test-by-test validity of the psychometric results based upon the Examiner's observations of the subject during testing and upon the Psychologist's evaluation of the data in test de-briefing with the subject. Use the numbered factors below to indicate the reason(s) for questionable validity among any of the data. For datum thought to be of questionable validity, also provide an estimate of the subject's "true" score or result. Forward the completed form with the subject's raw data.

Reasons for Questionable Validity

- Poor reading comprehension
 Fatigue
- Neg attitude, angry, marginal cooperator
- 4. Careless, hurried responses
- 5. Examiner Error

- Exaggeration of complaints ("fake bad")
- 7. Minimizing complaints ("fake good")
- 8. Disorganized personality (Psychotic)
- 9. Physically ill (flu, venipuncture effects, etc)
- 10. Other (Specify)

	<u>Test Score</u>	<u>Valid Results</u>	Reason(s) for Questionably Valid_Results	Est of "True" Score/Result
1.	WAIS VIQ PIQ FSIQ			
2. 3.	WRAT Reading Halstead-Reitan Category Test			

Ranch Hand II: Psychometric De-Briefing Form Continued

	Test_Score	Valid <u>Results</u>	Reason(s) for Questionably Valid Results	Est of "True" <u>Score/Result</u>
4.	Tactual Performance Test Preferred Hand Non-Preferred Hand Both Hands Memory Localization Speech-Sounds Perception Seashore Rhythm Finger Tapping Preferred Hand Non-Preferred Hand Trail Making Test Part A Part B Grip Strengths Preferred Hand Non-Preferred Hand WMS I Logical Mem (immed) Visual Repro (immed) Associate Lrng Logical Mem (delayed) Visual Repro (delayed) Visual Repro (delayed)	·		
5. 6.	MMPI (overall rating of protocol)			WNL or ONL

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