In the Listing of Impairments, the listings under each separate body system in both Part A and Part B will be effective for periods ranging from 4 to 8 years unless extended or revised and promulgated again. Specifically, the body system listings in the Listing of Impairments will be subject to the following termination dates:

- Musculoskeletal system (1.00) within 5 years. Consequently, the listings in this body system will no longer be effective on June 6, 1992.
- Respiratory system (3.00) within 6 years. Consequently, the listings in this body system will no longer be effective on December 6, 1991.
- The cardiovascular system (4.00) will no longer be effective on June 6, 1991.
- The listings under the other body systems in Part A and Part B will expire in 8 years. Consequently, the listing in these body systems will no longer be effective on December 6, 1993. The mental disorders listings in Part A will no longer be effective on August 28, 1991, unless extended by the Board or revised and promulgated again.

Part A

Criteria applicable to individuals age 18 and over and to children under age 18 where criteria are appropriate.

Sec.
1.00 Musculoskeletal System.
2.00 Special Senses and Speech.
3.00 Respiratory System.
4.00 Cardiovascular System.
5.00 Digestive System.
6.00 Genito-Urinary System.
7.00 Hemic and Lymphatic System.
8.00 Skin.
9.00 Endocrine System.
10.00 Multiple Body Systems.
11.00 Neurological.
12.00 Mental Disorders.
13.00 Neoplastic Diseases, Malignant.

1.00 Musculoskeletal System

A. Loss of function may be due to amputation or deformity. Pain may be an important factor in causing functional loss, but it must be demonstrated that it is a relevant abnormal sign or laboratory findings. Evaluations of musculoskeletal impairments should be supported where applicable by detailed descriptions of the joints, including ranges of motion, condition of the musculature, sensory or reflex changes, circulatory deficits, and X-ray abnormalities. (105C)

B. Disorders of the spine, associated with vertebrogenic disorders as in 1.05C, result in impairment because of distortion of the bony and ligamentous architecture of the spine or impingement of a herniated nucleus pulposus or bulging annulus on a nerve root. Impairment caused by such abnormalities usually improves with time or responds to treatment. (105C)

Criteria applicable to individuals age 18 and over and to children under age 18 where criteria are appropriate.

1.05C

The clinical diagnosis of the entity to be evaluated first must be established on the basis of adequate history, physical examination, and roentgenograms. The specific findings stated in 1.05C represent the level required for that impairment; these findings, by themselves, are not intended to represent the basis for establishing the clinical diagnosis. Furthermore, while neurological examination findings are required, they are not to be interpreted as a basis for evaluating the magnitude of any neurological impairment. Neurological impairments are to be evaluated under 11.00-11.19.

The history must include a detailed description of the character, location, and radiation of pain; mechanical factors which incite and relieve pain; prescribed treatment, including type, dose, and frequency of analgesic; and typical daily activities. Care must be taken to ascertain that the reported examination findings are consistent with the individual’s daily activities.

There must be a detailed description of the orthopedic and neurologic examination findings. The findings should include a description of gait, limitation of movement of the spine given quantitatively in degrees from the vertical position, motor and sensory abnormalities, muscle spasm, and deep tendon reflexes. Observations of the individual during the examination should be reported; e.g., how he or she gets on and off the examining table. Inability to walk on heels or toes, to squat, or to arise from a squatting position, where appropriate, may be considered evidence of significant motor loss. However, a report of atrophy is not acceptable as evidence of significant motor loss without circumferential measurements of both thighs and lower legs (or upper or lower arms) at a stated point above and below the knee or elbow given in inches or centimeters. A specific description of atrophy of hand muscles is acceptable without measurements of atrophy but should include measurements of grip strength.

These physical examination findings must be determined on the basis of objective observations during the examination and not simply a report of the individual’s allegation, e.g., he says his leg is week, numb, etc. Alternative testing methods should be used to verify the objectivity of the abnormal findings, e.g., a seated straight-leg raising.
test in addition to a supine straight-leg raising test. Since abnormal findings may be intermittent, their continuous presence over a period of time must be established by a record of ongoing treatment. Neurological abnormalities may not completely subside after surgical or nonsurgical treatment, or with the passage of time. Residual neurological abnormalities, which persist after it has been determined clinically or by direct surgical or other observation that the ongoing or progressive condition is no longer present, cannot be considered to satisfy the required findings in 1.05C.

Where surgical procedures have been performed, documentation should include a copy of the operative note and available pathology reports.

Electrodiagnostic procedures and myelography may be useful in establishing the clinical diagnosis, but do not constitute alternative criteria to the requirements in 1.05C.

C. After maximum benefit from surgical therapy has been achieved in situations involving fractures of an upper extremity (see 1.12) or soft tissue injuries of a lower or upper extremity (see 1.13), i.e., there have been no significant changes in physical findings or X-ray findings for any 6-month period after the last definitive surgical procedure, evaluation should be made on the basis of demonstrable residuals.

D. Major joints as used herein refer to hip, knee, ankle, shoulder, elbow, or wrist and hand. (Wrist and hand are considered together as one major joint.)

E. The measurements of joint motion are based on the techniques described in the "Joint Motion Method of Measuring and Recording," published by the American Academy of Orthopedic Surgeons in 1965, or the "Guides to the Evaluation of Permanent Impairment—The Extremities and Back" (Chapter I), American Medical Association, 1971.

1.01 Category of Impairments, Musculoskeletal

1.02 Active rheumatoid arthritis and other inflammatory arthritis.

With both A and B.

A. History of persistent joint pain, swelling, and tenderness involving multiple major joints (see 1.00D) and with signs of joint inflammation (swelling and tenderness) on current physical examination despite prescribed therapy for at least 3 months, resulting in significant restriction of function of the affected joints, and clinical activity expected to last at least 12 months; and

B. Corroboration of diagnosis at some point in time by either:

1. Positive serologic test for rheumatoid factor; or
2. Antinuclear antibodies; or
3. Elevated sedimentation rate; or
4. Characteristic histologic changes in biopsy of synovial membrane or subcutaneous nodule (obtained independent of Social Security disability evaluation).

1.03 Arthritis of a major weight-bearing joint (due to any cause):

With history of persistent joint pain and stiffness with signs of marked limitation of motion or abnormal motion of the affected joint on current physical examination. With:

A. Gross anatomical deformity of hip or knee (e.g., subluxation, contracture, bony or fibrous ankylosis, instability) supported by X-ray evidence of either significant joint space narrowing or significant bony destruction and markedly limiting ability to walk and stand; or

B. Reconstructive surgery or surgical arthrodesis of a major weight-bearing joint and return to full weight-bearing status did not occur, or is not expected to occur, within 12 months of onset.

1.04 Arthritis of one major joint in each of the upper extremities (due to any cause):

With history of persistent joint pain and stiffness, signs of marked limitation of motion of the affected joints on current physical examination, and X-ray evidence of either significant joint space narrowing or significant bony destruction. With:

A. Abduction and forward flexion (elevation) of both arms at the shoulders, including scapular motion, restricted to less than 90 degrees; or

B. Gross anatomical deformity (e.g., subluxation, contracture, bony or fibrous ankylosis, instability, ulnar deviation) and enlargement or effusion of the affected joints.

1.05 Disorders of the spine:

A. Arthritis manifested by ankylosis or fixation of the affected vertebral body at 30% or more of flexion measured from the neutral position, with X-ray evidence of:

1. Calcification of the anterior and lateral ligaments; or

B. Bilateral ankylosis of the sacroiliac joints with abnormal apophyseal articulations; or

C. Osteoporosis, generalized (established by X-ray) manifested by pain and limitation of back motion and paravertebral muscle spasm with X-ray evidence of:

1. Compression fracture of a vertebral body with loss of at least 50 percent of the estimated height of the vertebral body prior to the compression fracture, with no intervening direct traumatic episode; or

2. Multiple fractures of vertebrae with no intervening direct traumatic episode; or

C. Other vertebrogenic disorders (e.g., herniated nucleus pulposus, spinal stenosis) with the following persisting for at least 3 months despite prescribed therapy and expected to last 12 months. With both 1 and 2:

1. Pain, muscle spasm, and significant limitation of motion in the spine; and
Fractures of an upper extremity

1. Vascular disease; or
2. Neurological changes associated with vascular or neurological deficits, traumatic loss of muscle mass or tendons and X-ray evidence of bony ankylosis at an unfavorable angle, joint subluxation or instability:
   A. Both hands; or
   B. Both feet; or
   C. One hand and one foot.

1.09 Amputation or anatomical deformity of (i.e., loss of major function due to degenerative changes associated with vascular or neurological deficits, traumatic loss of muscle mass or tendons and X-ray evidence of bony ankylosis at an unfavorable angle, joint subluxation or instability):
   A. Hemipelvectomy or hip disarticulation; or
   B. Amputation at or above the tarsal region due to peripheral vascular disease or diabetes mellitus; or
   C. Inability to use a prosthesis effectively, without obligatory assistive devices, due to one of the following:
      1. Vascular disease; or
      2. Neurological complications (e.g., loss of position sense); or
      3. Stump too short or stump complications persistent, or are expected to persist, for at least 12 months from onset; or
      4. Disorder of contralateral lower extremity which markedly limits ability to walk and stand.

1.10 Amputation of one lower extremity (at or above the tarsal region):
   A. Non-union of a fracture of the shaft of the tibia, or a major joint of an upper or lower extremity, with persistent activity or occurrence of at least two episodes of acute activity or exacerbation of existing activity within a 5-month period prior to adjudication, manifested by local inflammatory, and systemic signs and laboratory findings (e.g., heat, redness, swelling, leucocytosis, or increased sedimentation rate) and expected to last at least 12 months despite prescribed therapy; or
   B. Multiple localizations and systemic manifestations as in A above.

1.11 Fracture of the femur, tibia, tarsal bone of pelvis with solid union not evident on X-ray and not clinically solid, when such determination is feasible, and return to full weight-bearing status did not occur or is not expected to occur within 12 months of onset.

1.12 Fractures of an upper extremity with non-union of a fracture of the shaft of the humerus, radius, or ulna under continuing surgical management directed toward restoration of functional use of the extremity and such function was not restored or expected to be restored within 12 months after onset.

1.13 Soft tissue injuries of an upper or lower extremity requiring a series of staged surgical procedures within 12 months after onset for salvage and/or restoration of major function of the extremity, and such major function was not restored or expected to be restored within 12 months after onset.

2. Appropriate radicular distribution of significant motor loss with muscle weakness and sensory and reflex loss.

1.08 Osteomyelitis or septic arthritis (established by X-ray):
   A. Located in the pelvis, vertebra, femur, tibia, or a major joint of an upper or lower extremity, with persistent activity or occurrence of at least two episodes of acute activity within a 5-month period prior to adjudication, manifested by local inflammatory, and systemic signs and laboratory findings (e.g., heat, redness, swelling, leucocytosis, or increased sedimentation rate) and expected to last at least 12 months despite prescribed therapy; or
   B. Amputation at or above the tarsal region due to peripheral vascular disease or diabetes mellitus; or
   C. Inability to use a prosthesis effectively, without obligatory assistive devices, due to one of the following:
      1. Vascular disease; or
      2. Neurological changes associated with vascular or neurological deficits, traumatic loss of muscle mass or tendons and X-ray evidence of bony ankylosis at an unfavorable angle, joint subluxation or instability:
         A. Hemipelvectomy or hip disarticulation; or
         B. Amputation at or above the tarsal region due to peripheral vascular disease or diabetes mellitus; or
         C. Inability to use a prosthesis effectively, without obligatory assistive devices, due to one of the following:
            1. Vascular disease; or
            2. Neurological complications (e.g., loss of position sense); or
            3. Stump too short or stump complications persistent, or are expected to persist, for at least 12 months from onset; or
            4. Disorder of contralateral lower extremity which markedly limits ability to walk and stand.

2.00 Special Senses and Speech

A. Ophthalmology

1. Causes of impairment. Diseases or injury of the eyes may produce loss of central or peripheral vision. Loss of central vision results in inability to distinguish detail and prevents reading and fine work. Loss of peripheral vision restricts the ability of an individual to move about freely. The extent of impairment of sight should be determined by visual testing.

2. Central visual acuity. A loss of central visual acuity may be caused by impaired distance vision. However, for an individual to meet the level of severity described in 2.02 and 2.04, only the remaining central visual acuity for distance of the better eye with best correction based on the Snellen test chart measurement may be used. Correction obtained by special visual aids (e.g., contact lenses) will be considered if the individual has the ability to wear such aids.

3. Field of vision. impairment of peripheral vision may result if there is contraction of the visual fields. The contraction may be either symmetrical or irregular. The extent of the remaining peripheral visual field will be determined by usual perimetric methods at a distance of 330 mm. under illumination of not less than 7-foot candles. For the phakic eye (the eye with a lens), a 3 mm. white disc target will be used, and for the aphakic eye (the eye without the lens), a 6 mm. white disc target will be used. In neither instance should corrective spectacle lenses be worn during the examination but if they have been used, this fact must be stated.

Measurements obtained on comparable perimetric devices may be used; this does not include the use of tangent screen measurements. For measurements obtained using the Goldmann perimeter, the object size designation III and the illumination designation 4 should be used for the phakic eye, and the object size designation IV and illumination designation 4 for the aphakic eye.

Field measurements must be accompanied by notated field charts, a description of the type and size of the target and the test distance. Tangent screen visual fields are not acceptable as a measurement of peripheral field loss.

Where the loss is predominately in the lower visual fields, a system such as the weighted grid scale for perimetric fields described by B. Esterman (see Grid for Scoring Visual Fields, II. Perimeter, Archives of Ophthalmology, 79:400, 1968) may be used for determining whether the visual field loss is comparable to that described in Table 2.

4. Muscle function. Paralysis of the third cranial nerve producing ptosis, paralysis of accommodation, and dilation and immobility of the pupil may cause significant visual
impaired. When all the muscle of the eye are paralyzed including the iris and ciliary body (total ophthalmoplegia), the condition is considered a severe impairment provided it is confirmed by a neuro-ophthalmologic examination which must be performed by or under the supervision of an ophthalmologist qualified to perform such tests.

Hearing tests should be preceded by an otolaryngologic examination with a detailed description of the vertiginous episodes, including notation of frequency, severity, and duration of the attacks. Pure tone and speech audiometry with the appropriate special examinations, such as Bekesy audiometry, are necessary. Vestibular functions are assessed by positional and caloric testing, preferably by electronystagmography. When polytograms, contrast radiography, or other special tests have been performed, copies of the reports of these tests should be submitted in addition to reports of skull and temporal bone X-rays.

3. Organic loss of speech. Glossectomy or laryngectomy or cicatricial laryngeal stenosis due to injury or infection results in

**Railroad Retirement Board**

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### Hearing Impairment

Speech discrimination should be determined using a standardized measure of speech discrimination ability in quiet at a test presentation level sufficient to ascertain maximum discrimination ability. The speech discrimination measure (test) used, and the level at which testing was done, must be reported.

Hearing tests should be preceded by an otolaryngologic examination with a detailed description of the vertiginous episodes, including notation of frequency, severity, and duration of the attacks. Pure tone and speech audiometry with the appropriate special examinations, such as Bekesy audiometry, are necessary. Vestibular functions are assessed by positional and caloric testing, preferably by electronystagmography. When polytograms, contrast radiography, or other special tests have been performed, copies of the reports of these tests should be submitted in addition to reports of skull and temporal bone X-rays.

3. Organic loss of speech. Glossectomy or laryngectomy or cicatricial laryngeal stenosis due to injury or infection results in
loss of voice production by normal means. In evaluating organic loss of speech (see 2.09), ability to produce speech by any means includes the use of mechanical or electronic devices. Impairment of speech due to neurologic disorders should be evaluated under 11.00-11.19.

2.01 Category of Impairments, Special Senses and Speech

2.02 Impairment of central visual acuity. Remaining vision in the better eye after best correction is 20/200 or less.

2.03 Contraction of peripheral visual fields in the better eye.
   A. To 10% or less from the point of fixation; or
   B. So the widest diameter subtends an angle no greater than 20°; or
   C. To 20 percent or less visual field efficiency.

2.04 Loss of visual efficiency. Visual efficiency of better eye after best correction 20 percent or less. (The percent of remaining central visual efficiency and the percent of remaining visual field efficiency.)

2.05 Complete homonymous hemianopsia (with or without macular sparing). Evaluate under 2.04.

2.06 Total bilateral ophthalmoplegia.

2.07 Disturbance of labyrinthine-vestibular function (including Meniere's disease), characterized by a history of frequent attacks of balance disturbance, tinnitus, and progressive loss of hearing. With both A and B:
   A. Disturbed function of vestibular labyrinth demonstrated by caloric or other vestibular tests; and
   B. Hearing loss established by audiometry.

2.08 Hearing impairments (hearing not restored by a hearing aid) manifested by:
   A. Average hearing threshold sensitivity for air conduction of 90 decibels or greater and for bone conduction to corresponding maximal levels, in the better ear, determined by the simple average of hearing threshold levels at 500, 1000 and 2000 Hz. (see 2.08 B1); or
   B. Speech discrimination scores of 40 percent or less in the better ear.

2.09 Organic loss of speech due to any cause with inability to produce by any means speech which can be heard understood and sustained.

1. Diagram of right eye illustrates extent of normal visual field as tested on standard perimeter at 3/330 (3 mm. white disc at a distance of 330 mm.) under 7 foot-candles illumination. The sum of the eight principal meridians of this field total 500°.

2. The percent of visual field efficiency is obtained by adding the number of degrees of the eight principal meridians of the contracted field and dividing by 500. Diagram of left eye illustrates visual field contracted to 30° in the temporal and down and out meridians and to 20° in the remaining six meridians. The percent of visual field efficiency of this field is: 6-20+2-30 =180/500=0.36 or 36 percent remaining visual field efficiency, or 64 percent loss.

### TABLE NO. 1—PERCENTAGE OF CENTRAL VISUAL EFFICIENCY CORRESPONDING TO CENTRAL VISUAL ACUITY NOTATIONS FOR DISTANCE IN THE PHAKIC AND APHAKIC EYE (BETTER EYE)

<table>
<thead>
<tr>
<th>Snellen</th>
<th>Percent central visual efficiency</th>
</tr>
</thead>
<tbody>
<tr>
<td>20/16</td>
<td>6/5</td>
</tr>
<tr>
<td>20/20</td>
<td>6/6</td>
</tr>
<tr>
<td>20/25</td>
<td>6/7.5</td>
</tr>
<tr>
<td>20/32</td>
<td>6/10</td>
</tr>
<tr>
<td>20/40</td>
<td>6/12</td>
</tr>
<tr>
<td>20/50</td>
<td>6/15</td>
</tr>
<tr>
<td>20/64</td>
<td>6/20</td>
</tr>
<tr>
<td>20/80</td>
<td>6/24</td>
</tr>
<tr>
<td>20/100</td>
<td>6/30</td>
</tr>
<tr>
<td>20/125</td>
<td>6/38</td>
</tr>
<tr>
<td>20/160</td>
<td>6/48</td>
</tr>
<tr>
<td>20/200</td>
<td>6/60</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>English</th>
<th>Metric</th>
<th>Phakic</th>
<th>Aphakic monocular</th>
<th>Aphakic binocular</th>
</tr>
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<tbody>
<tr>
<td>20/16</td>
<td>6/5</td>
<td>100</td>
<td>50</td>
<td>75</td>
</tr>
<tr>
<td>20/20</td>
<td>6/6</td>
<td>100</td>
<td>50</td>
<td>75</td>
</tr>
<tr>
<td>20/25</td>
<td>6/7.5</td>
<td>95</td>
<td>47</td>
<td>71</td>
</tr>
<tr>
<td>20/32</td>
<td>6/10</td>
<td>90</td>
<td>45</td>
<td>67</td>
</tr>
<tr>
<td>20/40</td>
<td>6/12</td>
<td>85</td>
<td>42</td>
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<tr>
<td>20/200</td>
<td>6/60</td>
<td>20</td>
<td></td>
<td></td>
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</tbody>
</table>

Column and Use.

1. Phakic.—1. A lens is present in both eyes. 2. A lens is present in the better eye and absent in the poorer eye. 3. A lens is present in one eye and the other eye is enucleated.

2. Monocular.—1. A lens is absent in the better eye and present in the poorer eye. 2. The lenses are absent in both eyes; however, the central visual acuity in the poorer eye after best correction is 20/200 or less. 3. A lens is absent from one eye and the other eye is enucleated.

3. Binocular.—1. The lenses are absent from both eyes and the central visual acuity in the poorer eye after best correction is greater than 20/200.
TABLE NO. 2—CHART OF VISUAL FIELD SHOWING EXTENT OF NORMAL FIELD AND METHOD OF COMPUTING PERCENT OF VISUAL FIELD EFFICIENCY

3.00 RESPIRATORY SYSTEM

A. Introduction: Impairments caused by the chronic disorder of the respiratory system generally result from irreversible loss of pulmonary functional capacity (ventilatory impairment, gas exchange impairment, or a combination of both). The most common symptom attributable to these disorders is dyspnea on exertion. Cough, wheezing, sputum production, hemoptysis, and chest pain may also occur, but need not be present. However, since these symptoms are common to many other diseases, evaluation of impairments of the respiratory system requires a history, physical examination, and chest roentgenogram to establish the diagnosis of a chronic respiratory disorder. Pulmonary function testing is required to provide a basis for assessing the impairment, once the diagnosis is established by appropriate clinical findings.

Alteration of ventilatory function may be due primarily to chronic obstructive pulmonary disease (emphysema, chronic bronchitis), chronic asthmatic bronchitis or restrictive disorders with primary loss of lung volume (pulmonary resection, thoracoplasty, chest cage deformity as seen in kyphoscoliosis), or infiltrative interstitial disorders (diffuse fibrosis). Impairment of gas exchange without significant airway obstruction may be produced by interstitial disorders (diffuse fibrosis). Primary disease of pulmonary circulation may produce pulmonary vascular hypertension and, eventually, heart failure. Whatever the mechanism, any chronic progressive pulmonary disorder may result in cor pulmonale or heart failure. Chronic infection caused, most frequently by mycobacterial or mycotic organisms, may produce extensive lung destruction resulting in marked loss of pulmonary functional capacity. Some disorders such as bronchiectasis and asthma may be characterized by acute, intermittent illnesses of such frequency and intensity that they produce a marked impairment apart from intercurrent functional loss, which may be mild.

Most chronic pulmonary disorders may be adequately evaluated on the basis of history, physical examination, chest roentgenogram, and ventilatory function tests. Direct assessment of gas exchange by exercise arterial blood gas determination or diffusing capacity is required only in specific relatively rare circumstances, depending on the clinical features and specific diagnosis.

B. Mycobacterial and mycotic infections of the lung will be evaluated on the basis of the resulting impairment to pulmonary function. Evidence of infectious or active mycobacterial or mycotic infection, such as positive cultures, increasing lesions, or cavitation, is not, by itself, a basis for determining that the individual has a severe impairment which is expected to last 12 months.
However, if these factors are abnormally persistent, they should not be ignored. For example, in those unusual cases where there is evidence of persistent pulmonary infection caused by mycobacterial or mycotic organisms for a period closely approaching 12 consecutive months, the clinical findings, complications, treatment considerations, and prognosis must be carefully assessed to determine whether, despite the absence of impairment of pulmonary function, the individual has a severe impairment that can be expected to last for 12 consecutive months.

C. When a respiratory impairment is episodic in nature, as may occur in complications of bronchiectasis and asthmatic bronchitis, the frequency of severe episodes despite prescribed treatment is the criterion for determining the level of impairment. Documentation for episodic asthma should include the hospital or emergency room records indicating the dates of treatment, clinical findings on presentation, what treatment was given and for what period of time, and the clinical response. Severe attacks of episodic asthma, as listed in section 3.03B, are defined as prolonged episodes lasting at least several hours, requiring intensive treatment such as intravenous drug administration or inhalation therapy in a hospital or emergency room.

D. Documentation of ventilatory function tests. The results of ventilatory function studies for evaluation under tables I and II should be expressed in liters or liters per minute (BTPS). The reported one second forced expiratory volume (FEV1) should be stated in the respiratory impairment meeting short of such purchase reveals that the impairment meets or equals any other listing or when the claim can be adjudicated on some other basis. Capillary blood analysis for PO2 or PCO2 is not acceptable. Analysis of arterial blood gases obtained after exercise is stopped is not acceptable.

Generally individuals with an FEV1 greater than 2.5 liters or an MVV greater than 100 liters per minute would not be considered for blood gas studies unless diffuse interstitial pulmonary fibrosis was noted on chest X-ray or documented by tissue diagnosis. The exercise test facility should be provided with the clinical reports, report of chest roentgenogram, and spirometry results obtained by the DDS. The testing facility should determine whether exercise testing is clinically contraindicated. If an exercise test is clinically contraindicated, the reason for exclusion from the test should be stated in the report of the exercise test facility.

2. Methodology. Individuals considered for exercise testing first should have resting PaO2, PaCO2, and pH determinations by the
testing facility. The samples should be obtained in the sitting or standing position. The individual should be exercised under steady state conditions, preferably on a treadmill for a period of 6 minutes at a speed and grade providing a workload of approximately 17 ml O₂/kg/min. If a bicycle ergometer is used, an exercise equivalent of 450 kpm/min., or 75 watts, should be used. At the option of the facility, a warm-up period of treadmill walking may be performed to acquaint the applicant with the procedure. If, during the warm-up period, the individual cannot exercise at the designated level, a lower speed and/or grade may be selected in keeping with the exercise capacity estimate. The individual should be monitored by electrocardiogram throughout the exercise and representative strips taken to provide heart rate in each minute of exercise. During the 5th or 6th minute of exercise, an arterial blood gas sample should be drawn and analyzed for PO₂, PCO₂, and pH. If the facility has the capability, and at the option of the DDS and the facility, minute ventilation (BTPS) and oxygen consumption per minute (STPD) and CO₂ production (STPD) should be measured during the 5th or 6th minute of exercise. If the individual fails to complete 6 minutes of exercise, the facility should comment on the reason.

The report should contain representative strips of electrocardiograms taken during the exercise, hematocrit, resting and exercise arterial blood gas value, speed and grade of the treadmill or bicycle ergometer exercise level in watts or kpm/min., and duration of exercise. The altitude of the test site, barometric pressure, and normal range of blood gas values for that facility should also be reported.

3. Evaluation. Three tables are provided in Listing 3.02C1 for evaluation of arterial blood gas determinations at rest and during exercise. The blood gas levels in Listing 3.02C1, Table III-A, are applicable at test sites situated at less than 3,000 feet above sea level. The blood gas levels in Listing 3.02C1, Table III-B, are applicable at test sites situated at 3,000 through 6,000 feet above sea level. The blood gas levels in Listing 3.02C1, Table III-C, are applicable for test sites over 6,000 feet above sea level. Tables III-B and C, take into account the lower blood PaO₂ normally found in individuals tested at the higher altitude. When the barometric pressure is unusually high for the altitude at the time of testing, consideration should be given to those cases in which the PaO₂ falls slightly above the requirements of Table III-A, III-B, or III-C, whichever is appropriate for the altitude at which testing was performed.

### TABLE I

<table>
<thead>
<tr>
<th>FEV₁ and MVV</th>
<th>VC equal to or less than (L, BTPS)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Height without shoes (inches)</td>
<td>(MBC) equal to or less than (L/min., BTPS)</td>
</tr>
<tr>
<td>60 or less</td>
<td>1.0</td>
</tr>
<tr>
<td>61-63</td>
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<tr>
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<td>66-67</td>
<td>1.3</td>
</tr>
<tr>
<td>68-69</td>
<td>1.4</td>
</tr>
<tr>
<td>70-71</td>
<td>1.5</td>
</tr>
<tr>
<td>72 or more</td>
<td>1.6</td>
</tr>
</tbody>
</table>

or

### TABLE II

<table>
<thead>
<tr>
<th>Height without shoes (inches)</th>
<th>VC equal to or less than (L, BTPS)</th>
</tr>
</thead>
<tbody>
<tr>
<td>60 or less</td>
<td>1.2</td>
</tr>
<tr>
<td>61-63</td>
<td>1.3</td>
</tr>
<tr>
<td>64-65</td>
<td>1.4</td>
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<tr>
<td>66-67</td>
<td>1.5</td>
</tr>
<tr>
<td>68-69</td>
<td>1.6</td>
</tr>
<tr>
<td>70-71</td>
<td>1.7</td>
</tr>
<tr>
<td>72 or more</td>
<td>1.8</td>
</tr>
</tbody>
</table>

or

### TABLE III-A

<table>
<thead>
<tr>
<th>Arterial PCO₂ (mm. Hg)</th>
<th>Arterial PO₂ and equal to or less than (mm. Hg)</th>
</tr>
</thead>
<tbody>
<tr>
<td>30 or below</td>
<td>65</td>
</tr>
<tr>
<td>31</td>
<td>64</td>
</tr>
<tr>
<td>32</td>
<td>63</td>
</tr>
<tr>
<td>33</td>
<td>62</td>
</tr>
</tbody>
</table>
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D. Mixed obstructive ventilatory and gas exchange impairment. Evaluate under the criteria in 3.02A, B, and C.

3.03 Asthma. With:
A. Chronic asthmatic bronchitis. Evaluate under the criteria for chronic obstructive ventilatory impairment in 3.02A, or
B. Episodes of severe attacks (See 3.00C), in spite of prescribed treatment, occurring at least once every 2 months or on an average of at least 6 times a year, and prolonged expiration with wheezing or rhonchi on physical examination between attacks.

3.06 Pneumoconiosis (demonstrated by roentgenographic evidence). Evaluate under criteria in 3.02.

3.07 Bronchiectasis (demonstrated by radiopaque material). With:
A. Episodes of acute bronchitis or pneumonia or hemoptysis (more than blood-streaked sputum) occurring at least every 2 months; or
B. Impairment of pulmonary function due to extensive disease should be evaluated under the applicable criteria in 3.02.

3.08 Mycobacterial infection of the lung. Impairment of pulmonary function due to extensive disease should be evaluated under appropriate criteria in 3.02.

3.09 Mycotic infection of the lung. Impairment of pulmonary function due to extensive disease should be evaluated under the appropriate criteria in 3.02.

3.11 Cor pulmonale, or pulmonary vascular hypertension. Evaluate under the criteria in 4.02.

4.00 Cardiovascular System

A. Severe cardiac impairment results from one or more of three consequences of heart disease; (1) congestive heart failure; (2) ischemia (with or without necrosis) of heart muscle; (3) conduction disturbances and/or arrhythmias resulting in cardiac syncope.

With diseases of arteries and veins, severe impairment may result from disorders of the vasculature in the central nervous system, eyes, kidneys, extremities, and other organs. The criteria for evaluating impairment resulting from heart diseases or diseases of the blood vessels are based on symptoms, physical signs and pertinent laboratory findings.

B. Congestive heart failure is considered in the Listing under one category whatever the etiology (i.e., arteriosclerotic, hypertensive, rheumatic, pulmonary, congenital, or other organic heart diseases). Congestive heart failure is not considered to have been established for the purpose of 4.02 unless there is evidence of vascular congestion such as hepatomegaly or peripheral or pulmonary edema which is consistent with clinical diagnosis. (Radiological description of vascular congestion, unless supported by appropriate clinical evidence, should not be construed as pulmonary edema.) The findings of vascular congestion need not be present at the time of

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**TABLE III—A**

(Applicable at test sites less than 3,000 feet above sea level)

<table>
<thead>
<tr>
<th>Arterial PO&lt;sub&gt;2&lt;/sub&gt; (mm. Hg)</th>
<th>Arterial PO&lt;sub&gt;2&lt;/sub&gt; and equal to or less than (mm. Hg)</th>
</tr>
</thead>
<tbody>
<tr>
<td>31</td>
<td>60</td>
</tr>
<tr>
<td>32</td>
<td>69</td>
</tr>
<tr>
<td>33</td>
<td>74</td>
</tr>
<tr>
<td>34</td>
<td>80</td>
</tr>
<tr>
<td>35</td>
<td>86</td>
</tr>
<tr>
<td>36</td>
<td>92</td>
</tr>
<tr>
<td>37</td>
<td>98</td>
</tr>
<tr>
<td>38</td>
<td>104</td>
</tr>
<tr>
<td>39</td>
<td>110</td>
</tr>
<tr>
<td>40 or above</td>
<td>116</td>
</tr>
</tbody>
</table>

**TABLE III—B**

(Applicable at test sites 3,000 through 6,000 feet above sea level)

<table>
<thead>
<tr>
<th>Arterial PO&lt;sub&gt;2&lt;/sub&gt; (mm. Hg)</th>
<th>Arterial PO&lt;sub&gt;2&lt;/sub&gt; and equal to or less than (mm. Hg)</th>
</tr>
</thead>
<tbody>
<tr>
<td>30 or below</td>
<td>60</td>
</tr>
<tr>
<td>31</td>
<td>66</td>
</tr>
<tr>
<td>32</td>
<td>72</td>
</tr>
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<td>33</td>
<td>78</td>
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<tr>
<td>34</td>
<td>84</td>
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<td>35</td>
<td>90</td>
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<td>36</td>
<td>96</td>
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<td>37</td>
<td>102</td>
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<tr>
<td>38</td>
<td>108</td>
</tr>
<tr>
<td>39</td>
<td>114</td>
</tr>
<tr>
<td>40 or above</td>
<td>120</td>
</tr>
</tbody>
</table>

**TABLE III—C**

(Applicable at test sites over 6,000 feet above sea level)

<table>
<thead>
<tr>
<th>Arterial PO&lt;sub&gt;2&lt;/sub&gt; (mm. Hg) and</th>
<th>Arterial PO&lt;sub&gt;2&lt;/sub&gt; and equal to or less than (mm. Hg)</th>
</tr>
</thead>
<tbody>
<tr>
<td>30 or below</td>
<td>56</td>
</tr>
<tr>
<td>31</td>
<td>62</td>
</tr>
<tr>
<td>32</td>
<td>68</td>
</tr>
<tr>
<td>33</td>
<td>74</td>
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<tr>
<td>34</td>
<td>80</td>
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<tr>
<td>35</td>
<td>86</td>
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<tr>
<td>36</td>
<td>92</td>
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<tr>
<td>37</td>
<td>98</td>
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<tr>
<td>38</td>
<td>104</td>
</tr>
<tr>
<td>39</td>
<td>110</td>
</tr>
<tr>
<td>40 or above</td>
<td>116</td>
</tr>
</tbody>
</table>

2. Diffusing capacity for the lungs for carbon monoxide less than 6 ml./mm. Hg/min. (steady-state methods) or less than 9 ml./mm. Hg/min. (single breath method) or less than 30 percent of predicted normal. (All method, actual values, and predicted normal values for the methods used should be reported.)
adjudication (except for 4.02A), but must be casually related to the current episode of marked impairment. The findings other than vascular congestion must be persistent.

G. Ischemic, hiatal hernia, peptic ulcer, and peptic ulcer, and pan- lesions such as biliary tract disease, esophagitis, hiatal hernia, peptic ulcer, and pan-

Chest pain that appears to be of cardiac origin may be caused by noncoronary conditions. Evidence for the latter should be actively considered in determining whether the chest pain is of cardiac origin. Among the more common conditions which may masquerade as angina are gastrointestinal tract lesions such as biliary tract disease, esophagitis, hiatal hernia, peptic ulcer, and pancreatic pain. Since the results of a treadmill exercise test are the primary basis for adjudicating claims under 4.04, they should be included in the file whenever they have been performed. There are also circumstances under which it will be appropriate to purchase exercise tests. Generally, these are limited to claims involving chest pain which is considered to be of cardiac origin but without corroborating ECG or other evidence of ischemic heart disease.

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I. Electrocardiograms obtained at rest must be submitted in the original or a legible copy of a 12-lead tracing appropriately labeled, with the standardization inscribed on the tracing. Alteration in standardization of specific leads (such as to accommodate large ORS amplitudes) must be shown on those leads.

The effect of drugs, electrolyte imbalance, etc., should be considered as possible noncoronary causes of ECG abnormalities, especially those involving the ST segment. If needed and available, pre-drug (especially predigitalis) tracing should be obtained.

The term “ischemic” is used in 4.04 to describe a pathologic ST deviation. Nonspecific repolarization changes should not be confused with ischemic configurations or a current of injury.

Detailed descriptions or computer interpretations without the original or legible copies of the ECG are not acceptable.

2. Electrocardiograms obtained in conjunction with exercise tests must include the original tracings or a legible copy of appropriate leads obtained before, during, and after exercise. Test control tracings, taken before exercise in the upright position, must be obtained. An ECG after 20 seconds of vigorous hyperventilation should be obtained. A posthyperventilation tracing may be essential for the proper evaluation of an “abnormal” test in certain circumstances, such as in women with evidence of mitral valve prolapse. A tracing should be taken at approximately 5 METs of exercise and at the time the ECG becomes abnormal according to the criteria in 4.04A. The time of onset of these abnormal changes must be noted, and the ECG tracing taken at the time should be obtained. Exercise histograms without the original tracings or legible copies are not acceptable.

Whenever electrocardiographically documented stress test data are submitted, irrespective of the type, the standardization must be inscribed on the tracings and the strips must be labeled appropriately, indicating the times recorded. The degree of exercise achieved, the blood pressure levels during the test, and any reason for terminating the test must be included in the report.

G. Exercise testing.

1. When to purchase. Since the results of a treadmill exercise test are the primary basis for adjudicating claims under 4.04, they should be included in the file whenever they have been performed. There are also circumstances under which it will be appropriate to purchase exercise tests. Generally, these are limited to claims involving chest pain which is considered to be of cardiac origin but without corroborating ECG or other evidence of ischemic heart disease.
Exercise test should not be purchased in the absence of alleged chest pain of cardiac origin. Even in the presence of an allegation of chest pain of cardiac origin, an exercise test should not be purchased where full development of such a purchase reveals that the impairment meets or equals any Listing or the claim can be adjudicated on some other basis.

2. Methodology. When an exercise test is purchased, it should be a treadmill type using a continuous progressive multistage regimen. The targeted heart rate should be not less than 85 percent of the maximum predicted heart rate unless it becomes hazardous to exercise to the heart rate or becomes unnecessary because the ECG meets the criteria in 4.04A at a lower heart rate (see also 4.0F.2). Beyond these requirements, it is prudent to accept the methodology of a qualified, competent test facility. In any case, a precise description of the protocol that was followed must be provided.

3. Limitations of exercise testing. Exercise testing should not be purchased for individuals who have the following: unstable progressive angina pectoris; recent onset (approximately 2 months) of angina; congestive heart failure; uncontrolled serious arrhythmias (including uncontrolled auricular fibrillation); second or third-degree heart block; Wolff-Parkinson-White syndrome; uncontrolled marked hypertension; marked aortic stenosis; marked pulmonary hypertension; dissecting or ventricular aneurysms; acute illness; limiting neurological or musculoskeletal impairments; or for individuals on medication where performance of stress testing may constitute a significant risk.

The presence of noncoronary or nonischemic factors which may influence the ECG response to exercise include hypokalemia, hypertentilation, vasoregulatory asthenia, significant anemia, left bundle branch block, and other heart disease, particularly valvular.

Digitalis may cause ST segment abnormalities at rest, during, and after exercise. Digitalis-related ST depression, present at rest, may become accentuated and result in false interpretations of the ECG taken during or after exercise test.

4. Evaluation. Where the evidence includes the results of a treadmill exercise test, this evidence is the primary basis for adjudicating claims under 4.04. For purposes of this Social Security disability program, treadmill exercise testing will be evaluated on the basis of the level at which the test becomes positive in accordance with the ECG criteria in §404A. However, the significance of findings of a treadmill exercise test must be considered in light of the clinical course of the disease which may have occurred subsequent to performance of the exercise test. The criteria in 4.04B are not applicable if there is documentation of an acceptable treadmill exercise test, if there is no evidence of a treadmill exercise test or if the test is not acceptable, the criteria in 4.04B should be used. The level of exercise is considered in terms of multiples of MET’s (metabolic equivalent units). One MET is the basal O2 requirement of the body in an inactive state, sitting quietly. It is considered by most authorities to be approximately 3.5 ml. O2/kg/min.

H. Angiographic evidence.

1. Coronary arteriography. This procedure is not to be purchased by the Social Security Administration. Should the results of such testing be available, the report should be considered as to the quality and kind of data provided and its applicability to the requirements of the Listing of Impairments. A copy of the report of the catheterization and ancillary studies should be obtained. The report should provide information as to the technique used, the method of assessing coronary lumen diameter, and the nature and location of any obstructive lesions.

It is helpful to know the method used, the number of projections, and whether selective engagement of each coronary vessel was satisfactorily accomplished. It is also important to know whether the injected vessel was entirely and uniformly opacified, thus avoiding the artificial appearance of narrowing or an obstruction.

Coronary artery spasm induced by intracoronary catheterization is not to be considered as evidence of ischemic heart disease.

Estimation of the functional significance of an obstructive lesion may also be aided by description of how well the distal part of the vessel is visualized. Some patients with significant proximal coronary atherosclerosis have well-developed large collateral blood supply to the distal vessels without evidence of myocardial damage or ischemia, even under conditions of severe stress.

2. Left ventriculography. The report should describe the local contractility of the myocardium as may be evident from areas of hypokinesia, dyskinesia, or akinesia; and the overall contractility of the myocardium as measured by the ejection fraction.

3. Proximal coronary arteries (see 4.0B.7) will be considered as the:
   a. Right coronary artery proximal to the acute marginal branch; or
   b. Left anterior descending coronary artery proximal to the first septal perforator; or
   c. Left circumflex coronary artery proximal to the first obtuse marginal branch.

I. Results of other tests. Information from adequate reports of other tests such as radionuclide studies or echocardiography should be considered where that information is comparable to the requirements in the listing. An ejection fraction measured by echocardiography is not determinative, but may
be given consideration in the context of associated findings.

J. Major surgical procedures. The amount of function restored and the time required to effect an improvement after heart or vascular surgery vary with the nature and extent of the disorder, the type of surgery, and other individual factors. If the criteria described for heart or vascular disease are met, proposed heart or vascular surgery (coronary artery bypass procedure, valve replacement, major arterial grafts, etc.) does not militate against a finding of disability with subsequent assessment postoperatively.

The usual time after surgery for adequate assessment of the results of surgery is considered to be approximately 3 months. Assessment of the magnitude of the impairment following surgery requires adequate documentation of the pertinent evaluations and tests performed following surgery, such as an interval history and physical examination, with emphasis on those signs and symptoms which might have changed postoperatively, as well as X-rays and electrocardiograms. Where treadmill exercise tests or angiography have been performed following the surgical procedure, the results of these tests should be obtained.

Documentation of the preoperative evaluation and a description of the surgical procedure are also required. The evidence should be documented from hospital records (catheterization reports, coronary arteriographic reports, etc.) and the operative note.

Implantation of a cardiac pacemaker is not considered a major surgical procedure for purposes of this section.

K. Evaluation of peripheral arterial disease. The evaluation of peripheral arterial disease is based on medically acceptable clinical findings providing adequate history and physical examination findings describing the impairment, and on documentation of the appropriate laboratory techniques. The specific findings stated in Listing 4.13 represent the level of severity of that impairment; these findings, by themselves, are not intended to represent the basis for establishing the clinical diagnosis. The level of the impairment is based on the symptomatology, physical findings, Doppler studies before and after a standard exercise test, and/or angiographic findings.

The requirements for evaluation of peripheral arterial disease in Listing 4.13B are based on the ratio of systolic blood pressure at the ankle and other pertinent levels measured after exercise, and the time required to return the systolic blood pressure to the preexercise level. When exercise Doppler studies are purchased by the Social Security Administration, it is suggested that the requested exercise be on a treadmill at 2 mph, on a 12 percent grade for 5 minutes. Exercise studies should not be performed on individuals for whom exercise is contraindicated. The methodology of a qualified, competent facility should be accepted. In any case, a precise description of the protocol that was followed must be provided.

It must be recognized that application of the criteria in Listing 4.13B may be limited in individuals who have severe calcific (Monckeberg’s) sclerosis of the peripheral arteries or severe small vessel disease in individuals with diabetes mellitus.

Listing 4.13B.1 provides for determining that the listing is met when the resting anklebrachial systolic blood pressure ratio is less than 0.50. Listing 4.13B.2 provides additional criteria for evaluating peripheral arterial impairment on the basis of exercise studies when the resting anklebrachial systolic blood pressure ratio is 0.50 or above. The results of exercise studies should describe the level of exercise (e.g., speed and grade of the treadmill settings), the duration of exercise, symptoms during exercise, the reasons for stopping exercise if the expected level of exercise was not attained, blood pressures at the ankle and other pertinent levels measured after exercise, and the time required to return the systolic blood pressure to the preexercise level.

When exercise Doppler studies are purchased by the Social Security Administration, it is suggested that the requested exercise be on a treadmill at 2 mph, on a 12 percent grade for 5 minutes. Exercise studies should not be performed on individuals for whom exercise is contraindicated. The methodology of a qualified, competent facility should be accepted. In any case, a precise description of the protocol that was followed must be provided.

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Listing 4.13B.1 provides for determining that the listing is met when the resting anklebrachial systolic blood pressure ratio is less than 0.50. Listing 4.13B.2 provides additional criteria for evaluating peripheral arterial impairment on the basis of exercise studies when the resting anklebrachial systolic blood pressure ratio is 0.50 or above. The results of exercise studies should describe the level of exercise (e.g., speed and grade of the treadmill settings), the duration of exercise, symptoms during exercise, the reasons for stopping exercise if the expected level of exercise was not attained, blood pressures at the ankle and other pertinent levels measured after exercise, and the time required to return the systolic blood pressure to the preexercise level.

When exercise Doppler studies are purchased by the Social Security Administration, it is suggested that the requested exercise be on a treadmill at 2 mph, on a 12 percent grade for 5 minutes. Exercise studies should not be performed on individuals for whom exercise is contraindicated. The methodology of a qualified, competent facility should be accepted. In any case, a precise description of the protocol that was followed must be provided.

It must be recognized that application of the criteria in Listing 4.13B may be limited in individuals who have severe calcific (Monckeberg’s) sclerosis of the peripheral arteries or severe small vessel disease in individuals with diabetes mellitus.

Listing 4.13B.1 provides for determining that the listing is met when the resting anklebrachial systolic blood pressure ratio is less than 0.50. Listing 4.13B.2 provides additional criteria for evaluating peripheral arterial impairment on the basis of exercise studies when the resting anklebrachial systolic blood pressure ratio is 0.50 or above. The results of exercise studies should describe the level of exercise (e.g., speed and grade of the treadmill settings), the duration of exercise, symptoms during exercise, the reasons for stopping exercise if the expected level of exercise was not attained, blood pressures at the ankle and other pertinent levels measured after exercise, and the time required to return the systolic blood pressure to the preexercise level.

When exercise Doppler studies are purchased by the Social Security Administration, it is suggested that the requested exercise be on a treadmill at 2 mph, on a 12 percent grade for 5 minutes. Exercise studies should not be performed on individuals for whom exercise is contraindicated. The methodology of a qualified, competent facility should be accepted. In any case, a precise description of the protocol that was followed must be provided.

It must be recognized that application of the criteria in Listing 4.13B may be limited in individuals who have severe calcific (Monckeberg’s) sclerosis of the peripheral arteries or severe small vessel disease in individuals with diabetes mellitus.
D. Cor pulmonale (non-acute) documented by:
1. Right ventricular enlargement (or prominence of the right out-flow tract) on chest roentgenogram or fluoroscopy; and
2. ECG evidence of right ventricular hypertrophy with R wave of 5.0 mm. or greater in lead V1 and progressive decrease in R/S amplitude from lead V1 to V6.

4.03 Hypertensive vascular disease. Evaluate under 4.02 4.04 or under the criteria for the affected body system.

4.04 Ischemic heart disease with chest pain or cardiac origin as described in 4.00E With:
A. Treadmill exercise test (see 4.00 F and (G) demonstrating one of the following at an exercise level of 5 METS or less:
   1. Horizontal or downsloping depression (from the standing control) of the ST segment to 1.0 mm. or greater, lasting for at least 0.08 second after the J junction, and clearly discernible in at least two consecutive complexes which are on a level baseline in any lead; or
   2. J uncional depression occurring during exercise, remaining depressed (from the standing control) to 2.0 mm. or greater for at least 0.08 second after the J junction (the so-called slow upsloping ST segment), and clearly discernible in at least two consecutive complexes which are on a level baseline in any lead; or
   3. Premature ventricular systoles which are multiform or bidirectional or are sequentially inscribed (3 or more); or
   4. ST segment elevation (from the standing control) to 1 mm. or greater; or
   5. Development of second or third degree heart block; or
B. In the absence of a report of an acceptable treadmill exercise test (see 4.00G), one of the following:
   1. Transmural myocardial infarction exhibiting a Q5 pattern or a Q wave with amplitude at least 3rd of R wave and with a duration of 0.04 second or more. (If these are present in leads III and a VF: only, the requisite Q wave findings must be shown, by labelled tracing, to persist on deep inspiration); or
   2. Resting ECG findings showing ischemic-type (see 4.00F 1) depression of ST segment to more than 0.5 mm. in either (a) leads I and a VL and V6 or (b) leads II and III and aVF or (c) leads V1 through V6; or
   3. Resting ECG findings showing an ischemic configuration or current of injury (see 4.00F 1) with ST segment elevation to 2 mm. or more in either (a) leads I and a VL and V6 or (b) leads II and III and aVF or (c) leads V1 through V6; or
   4. Resting ECG findings showing symmetrical inversion of T waves to 5.0 mm. or more in any two leads except leads III or aVR or V1 or V6; or
   5. Inversion of T wave to 1.0 mm. or more in any of leads I, II, aVL, V2 to V6 and R wave of 5.0 mm. or more in lead aVL and R wave greater than S wave in lead aVF; or
   6. “Double” Master Two-Step test demonstrating one of the following:
      a. Ischemic depression of ST segment to more than 0.5 mm. lasting for at least 0.08 second beyond the J junction and clearly discernible in at least two consecutive complexes which are on a level baseline in any lead; or
      b. Development of a second or third degree heart block; or
   7. Angiographic evidence (see 4.00H) (obtained independent of Social Security disability evaluation) showing one of the following:
      a. 50 percent or more narrowing of the left main coronary artery; or
      b. 70 percent or more narrowing of a proximal coronary artery (see 4.00H3) (excluding the left main coronary artery); or
      c. 50 percent or more narrowing involving a long (greater than 1 cm.) segment of a proximal coronary artery or multiple proximal coronary arteries; or
   8. Akinetic or hypokinetic myocardial wall or septal motion with left ventricular ejection fraction of 30 percent of less measured by contrast or radio-isotopic ventriculographic methods; or
   C. Resting ECG findings showing left bundle branch block as evidenced by QRS duration of 0.12 second or more in leads I, II, or III and R peak duration of 0.06 second or more in leads I, aVL, V5, or V6, unless there is a coronary angiogram of record which is negative (see criteria in 4.04B7).

4.05 Recurrent arrhythmias (not due to digi-talis toxicity) resulting in uncontrolled repeated episodes of cardiac syncope and documented by resting or ambulatory (Holter) electrocardiography.

4.09 Myocardiopathies, rheumatic or syphilitic heart disease. Evaluate under the criteria in 4.02, 4.04, 4.05, or 11.04.

4.11 Aneurysm of aorta or major branches (demonstrated by roentgenographic evidence). With:
A. Acute or chronic dissection not controlled by prescribed medical or surgical treatment; or
B. Congestive heart failure as described under the criteria in 4.02; or
C. Renal failure as described under the criteria in 6.02; or
D. Repeated sncopal episodes.

4.12 Chronic venous insufficiency of the lower extremity with incompetency or obstruction of the deep venous return, associated with superficial varicosities, extensive brawny edema, stasis dermatitis, and recurrent or persistent ulceration which has not healed following at least 3 months of prescribed medical or surgical therapy.

4.13 Peripheral arterial disease. With:
A. Intermittent claudication with failure to visualize (on arteriogram obtained independent of Social Security disability evaluation) the common femoral or deep femoral artery in one extremity; or
B. Intermittent claudication with marked impairment of peripheral arterial circulation as determined by Doppler studies showing:
   1. Resting anklebrachial systolic blood pressure ratio of less than 0.50; or
   2. Decrease in systolic blood pressure at ankle or exercise (see 4.00K) to 50 percent or more of preexercise level and requiring 10 minutes or more to return to preexercise level; or
   C. Amputation at or above the tarsal region due to peripheral arterial disease.

5.00 DIGESTIVE SYSTEM

A. Disorders of the digestive system which result in a marked impairment usually do so because of interference with nutrition, multiple recurrent inflammatory lesions, or complications of disease, such as fistulae, abscesses, or recurrent obstruction. Such complications usually respond to treatment. These complications must be shown to persist on repeated examinations despite therapy for a reasonable presumption to be made that a marked impairment will last for a continuous period of at least 12 months.

B. Malnutrition or weight loss from gastrointestinal disorders. When the primary disorder of the digestive tract has been established (e.g., enterocolitis, chronic pancreatitis, postgastrointestinal resection, or esophageal stricture, stenosis, or obstruction), the resultant interference with nutrition will be considered under the criteria in 5.08. This will apply whether the weight loss is due to primary or secondary disorders of malabsorption, malassimilation or obstruction. However, weight loss not due to diseases of the digestive tract, but associated with psychiatric or primary endocrine or other disorders, should be evaluated under the appropriate criteria for the underlying disorder.

C. Surgical diversion of the intestinal tract, including colostomy or ileostomy, are not listed since they do not represent impairments which preclude all work activity if the individual is able to maintain adequate nutrition and function of the stoma. Dumping syndrome which may follow gastric resection rarely represents a marked impairment which would continue for 12 months. Peptic ulcer disease with recurrent ulceration after definitive surgery ordinarily responds to treatment. A recurrent ulcer after definitive surgery must be demonstrated on repeated upper gastrointestinal roentgenograms or gastroscopic examinations despite therapy to be considered a severe impairment which will last for at least 12 months. Definitive surgical procedures are those designed to control the ulcer disease process (i.e., vagotomy and pyloroplasty, subtotal gastrectomy, etc.). Simple closure of a perforated ulcer does not constitute definitive surgical therapy for peptic ulcer disease.

5.01 Category of Impairments, Digestive System

5.02 Recurrent upper gastrointestinal hemorrhage from undetermined cause with anemia manifested by hematocrit of 30 percent or less on repeated examinations.

5.03 Stricture, stenosis, or obstruction of the esophagus (demonstrated by X-ray or endoscopy) with weight loss as described under §5.08.

5.04 Peptic ulcer disease (demonstrated by X-ray or endoscopy). With:
   A. Recurrent ulceration after definitive surgery persistent despite therapy; or
   B. Inoperable fistula formation; or
   C. Recurrent obstruction demonstrated by X-ray or endoscopy, or
   D. Weight loss as described under §5.08.

5.05 Chronic liver disease (e.g., portal, postnecrotic, or biliary cirrhosis; chronic active hepatitis; Wilson’s disease). With:
   A. Esophageal varices (demonstrated by X-ray or endoscopy) with a documented history of massive hemorrhage attributable to these varices. Consider under a disability for 3 years following the last massive hemorrhage; thereafter, evaluate the residual impairment; or
   B. Performance of a shunt operation for esophageal varices. Consider under a disability for 3 years following surgery; thereafter, evaluate the residual impairment; or
   C. Ascites, not attributable to other causes, recurrent or persisting for at least 5 months, demonstrated by abdominal paracentesis or liver biopsy (obtained independent of Social Security disability evaluation) and one of the following:
      1. Ascites not attributable to other causes, recurrent or persisting for at least 3 months, demonstrated by abdominal paracentesis or associated with persistent hypoalbuminemia of 3.0 gm. per deciliter (100 ml.) or less; or
      2. Serum bilirubin of 2.5 mg. per deciliter (100 ml.) or greater on repeated examinations for at least 5 months; or
      D. Ascites, not attributable to other causes, recurrent or persisting for at least 5 months, demonstrated by abdominal paracentesis or associated with persistent hypoalbuminemia of 3.0 gm. per deciliter (100 ml.) or less; or
   E. Hepatic encephalopathy. Evaluate under the criteria in listing 12.02; or
   F. Confirmation of chronic liver disease by liver biopsy (obtained independent of Social Security disability evaluation) and one of the following:
      1. Ascites not attributable to other causes, recurrent or persisting for at least 3 months, demonstrated by abdominal paracentesis or associated with persistent hypoalbuminemia of 3.0 gm. per deciliter (100 ml.) or less; or
      2. Serum bilirubin of 2.5 mg. per deciliter (100 ml.) or greater on repeated examinations for at least 3 months; or
      3. Hepatic cell necrosis or inflammation, persisting for at least 3 months, documented by repeated abnormalities of prothrombin time and enzymes indicative of hepatic dysfunction.

5.06 Chronic ulcerative or granulomatous colitis (demonstrated by endoscopy, barium enema, biopsy, or operative findings). With:
A. Recurrent bloody stools documented on repeated examinations and anemia manifested by hematocrit of 30 percent or less on repeated examinations; or

B. Persistent or recurrent systemic manifestations, such as arthritis, iritis, fever, or liver dysfunction, not attributable to other causes; or

C. Intermittent obstruction due to intractable abscess, fistula formation, or stenosis; or

D. Recurrence of findings of A, B, or C above after total colectomy; or

E. Weight loss as described under §5.08.

5.07 Regional enteritis (demonstrated by operative findings, barium studies, biopsy, or endoscopy). With:

A. Persistent or recurrent intestinal obstruction evidenced by abdominal pain, distention, nausea, and vomiting and accompanied by stenotic areas of small bowel with proximal intestinal dilation; or

B. Persistent or recurrent systemic manifestations such as arthritis, iritis, fever, or liver dysfunction, not attributable to other causes; or

C. Intermittent obstruction due to intractable abscess or fistula formation; or

D. Weight loss as described under §5.08.

5.08 Weight loss due to any persisting gastrointestinal disorder (the following weights are to be demonstrated to have persisted for at least 3 months despite prescribed therapy and expected to persist at this level for at least 12 months.) With:

A. Weight equal to or less than the values specified in Table I or II; or

B. Weight equal to or less than the values specified in Table III or IV and one of the following abnormal findings on repeated examinations:

1. Serum albumin of 3.0 gm. per deciliter (100 ml.) or less; or

2. Hematocrit of 30 percent or less; or

3. Serum calcium of 8.0 mg. per deciliter (100 ml.) (4.0 mEq./L) or less; or

4. Uncontrolled diabetes mellitus due to pancreatic dysfunction with repeated hyperglycemia, hypoglycemia, or ketosis; or

5. Fat in stool of 7 gm. or greater per 24-hour stool specimen; or

6. Nitrogen in stool of 3 gm. or greater per 24-hour specimen; or

7. Persistent or recurrent ascites or edema not attributable to other causes.

Tables of weight reflecting malnutrition scaled according to height and sex—To be used only in connection with §5.08.

### Table I—Men—Continued

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1 Height measured without shoes.

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1 Height measured without shoes.

### Table III—Men

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1 Height measured without shoes.

### Table IV—Women

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6.02 Impairment of renal function, due to any chronic renal disease expected to last 12 months (e.g., hypertensive vascular disease, chronic nephritis, nephrolithiasis, polycystic disease, bilateral hydronephrosis, etc.) With:
A. Chronic hemodialysis or peritoneal dialysis necessitated by irreversible renal failure; or
B. Kidney transplant. Consider under a disability for 12 months following surgery; thereafter, evaluate the residual impairment (see 6.06C); or
C. Persistent elevation of serum creatinine in to 4 mg. per deciliter (100 ml.) or greater or reduction of creatinine clearance to 20 ml. per minute (29 liters/24 hours) or less, over at least 3 months, with one of the following:
1. Renal osteodystrophy manifested by severe bone pain and appropriate radiographic abnormalities (e.g., ostetis fibrosa, marked osteoporosis, pathologic fractures); or
2. A clinical episode of pericarditis; or
3. Persistent motor or sensory neuropathy; or
4. Intractable pruritus; or
5. Persistent fluid overload syndrome resulting in diastolic hypertension (110 mm. or above) or signs of vascular congestion; or
6. Persistent anorexia with recent weight loss and current weight meeting the values in 5.08, Table III or IV; or
7. Persistent hematocrits of 30 percent or less.
6.06 Nephrotic syndrome, with significant anasarca, persistent for at least 3 months despite prescribed therapy. With:
A. Serum albumin of 3.0 gm. per deciliter (100 ml.) or less and proteinuria of 3.5 gm. per 24 hours or greater;
B. Proteinuria of 10.0 gm. per 24 hours or greater.

7.00 HEMIC AND LYMPHATIC SYSTEM
A. Impairment caused by anemia should be evaluated according to the ability of the individual to adjust to the reduced oxygen carrying capacity of the blood. A gradual reduction in red cell mass, even to very low values, is often well tolerated in individuals with a healthy cardiovascular system.
B. Chronicity is indicated by persistence of the condition for at least 3 months. The laboratory findings cited must reflect the values reported on more than one examination over that 3-month period.
C. Sickle cell disease refers to a chronic hemolytic anemia associated with sickle cell hemoglobin, either homozygous or in combination with thalassemia or with another abnormal hemoglobin (such as C or F).

Appropriate hematologic evidence for sickle cell disease, such as hemoglobin electrophoresis, must be included. Vasocclusive or aplastic episodes should be documented by description of severity, frequency, and duration.
Major visceral episodes include meningitis, osteomyelitis, pulmonary infections or infarctions, cerebrovascular accidents, congestive heart failure, genito-urinary involvement, etc.

D. Coagulation defects. Chronic inherited coagulation disorders must be documented by appropriate laboratory evidence. Prophylactic therapy such as with antihemophilic globulin (AHG) concentrate does not in itself imply severity.

E. Acute leukemia. Initial diagnosis of acute leukemia must be based upon definitive bone marrow pathologic evidence. Recurrent disease may be documented by peripheral blood, bone marrow, or cerebrospinal fluid examination. The pathology report must be included.

The acute phase of chronic myelocytic (granulocytic) leukemia should be considered under the requirements for acute leukemia.

The criteria in 7.11 contain the designated duration of disability implicit in the finding of a listed impairment. Following the designated time period, a documented diagnosis itself is no longer sufficient to establish a marked impairment. The level of any remaining impairment must be evaluated on the basis of the medical evidence.

7.01 Category of Impairments, Hemic and Lymphatic System

7.02 Chronic anemia (hematocrit persisting at 30 percent or less due to any cause). With:

A. Requirement of one or more blood transfusions on an average of at least once every 2 months; or

B. Evaluation of the resulting impairment under criteria for the affected body system.

7.05 Sickle cell disease, or one of its variants. With:

A. Documented painful (thrombotic) crises occurring at least three times during the 5 months prior to adjudication; or

B. Requiring extended hospitalization (beyond emergency care) at least three times during the 12 months prior to adjudication; or

C. Chronic, severe anemia with persistence of hematocrit of 26 percent or less; or

D. Evaluate the resulting impairment under the criteria for the affected body system.

7.06 Chronic thrombocytopenia (due to any cause) with platelet counts repeatedly below 40,000/cubic millimeter. With:

A. At least one spontaneous hemorrhage, requiring transfusion, within 5 months prior to adjudication; or

B. Intracranial bleeding within 12 months prior to adjudication.

7.07 Hereditary telangiectasia with hemorrhage requiring transfusion at least three times during the 5 months prior to adjudication.

7.08 Coagulation defects (hemophilia or a similar disorder) with spontaneous hemorrhage requiring transfusion at least three times during the 5 months prior to adjudication.

7.09 Polycythemia vera (with erythrocytosis, splenomegaly, and leukocytosis or thrombocytosis). Evaluate the resulting impairment under the criteria for the affected body system.

7.10 Myelofibrosis (myeloproliferative syndrome). With:

A. Chronic anemia. Evaluate according to the criteria of §7.02; or

B. Documented recurrent systemic bacterial infections occurring at least 3 times during the 5 months prior to adjudication; or

C. Intractable bone pain with radiologic evidence of osteosclerosis.

7.11 Acute leukemia. Consider under a disability for 2½ years from the time of initial diagnosis.

7.12 Chronic leukemia. Evaluate according to the criteria of 7.02, 7.06, 7.10B, 7.11, 7.17, or 13.06A.

7.13 Lymphomas. Evaluate under the criteria in 13.06A.

7.14 Macroglobulinemia or heavy chain disease, confirmed by serum or urine protein electrophoresis or immunoelectrophoresis. Evaluate impairment under criteria for affected body system or under 7.02, 7.06, or 7.08.

7.15 Chronic polycythemia (due to any cause). With both A and B:

A. Absolute neutrophil counts repeatedly below 1,000 cells/cubic millimeter; and

B. Documented recurrent systemic bacterial infections occurring at least 3 times during the 5 months prior to adjudication.

7.16 Myeloma (confirmed by appropriate serum or urine protein electrophoresis and bone marrow findings). With:

A. Radiologic evidence of bony involvement with intractable bone pain; or

B. Evidence of renal impairment as described in 6.02; or

C. Hypercalcemia with serum calcium levels persistently greater than 11 mg. per deciliter (100 ml.) for at least 1 month despite prescribed therapy; or

D. Plasma cells (100 or more cells/cubic millimeter) in the peripheral blood.

7.17 Aplastic anemias or hematologic malignancies (excluding acute leukemia): With bone marrow transplantation. Consider under a disability for 12 months following transplantation; thereafter, evaluate according to the primary characteristics of the residual impairment.

8.00 SKIN

A. Skin lesions may result in a marked, long-lasting impairment if they involve extensive body areas or critical areas such as the hands or feet and become resistant to treatment. These lesions must be shown to have persisted for a sufficient period of time despite therapy for a reasonable presumption to be made that a marked impairment will
9.00 **Endocrine System**

Cause of impairment. Impairment is caused by overproduction or underproduction of hormones, resulting in structural or functional changes in the body. Where involvement of other organ systems has occurred as a result of a primary endocrine disorder, these impairments should be evaluated according to the criteria in the appropriate sections.

9.01 **Category of Impairments, Endocrine**

9.02 Thyroid Disorders. With:

A. Progressive exophthalmos as measured by exophthalmometry; or
B. Evaluate the resulting impairment under the criteria for the affected body system.

9.03 Hyperparathyroidism. With:

A. Generalized decalcification of bone on X-ray study and elevation of plasma calcium to 11 mg. per deciliter (100 ml.) or greater; or
B. A resulting impairment. Evaluate according to the criteria in the affected body system.

9.04 Hypoparathyroidism. With:

A. Severe recurrent tetany; or
B. Recurrent generalized convulsions; or
C. Lenticular cataracts. Evaluate under the criteria in 2.00ff.

9.05 Neurohypophyseal insufficiency (diabetes insipidus). With urine specific gravity of 1.005 or below, persistent for at least 3 months and recurrent dehydration.

9.06 Hyperfunction of the adrenal cortex. Evaluate the resulting impairment under the criteria for the affected body system.

9.08 Diabetes mellitus. With:

A. Neuropathy demonstrated by significant and persistent disorganization of motor function in two extremities resulting in sustained disturbance of gross and dexterous movements, or gait and station (see 11.00C); or
B. Acidosis occurring at least on the average of once every 2 months documented by appropriate blood chemical tests (pH or pCO2 or bicarbonate levels); or
C. Amputation at, or above, the tarsal region due to diabetic necrosis or peripheral arterial disease; or
D. Retinitis proliferans; evaluate the visual impairment under the criteria in 2.02, 2.03, or 2.04.

10.00 **Multiple Body Systems**

10.01 **Category of Impairments, Multiple Body Systems**

10.02 Hansen’s disease (leprosy). As active disease or consider as “under a disability” while hospitalized.

10.03 Polyarteritis or periarteritis nodosa (established by biopsy). With signs of generalized arterial involvement.

10.04 Disseminated lupus erythematosus (established by a positive LE preparation or biopsy or positive ANA test). With frequent exacerbations demonstrating involvement of renal or cardiac or pulmonary or gastrointestinal or central nervous systems.

10.05 Scleroderma or progressive systemic sclerosis (the diffuse or generalized form). With:

A. Advanced limitation of use of hands due to sclerodactyly or limitation in other joints; or
B. Significant visceral manifestations of digestive, cardiac, or pulmonary impairment.

10.10 Obesity. Weight equal to or greater than the values specified in Table I for
males. Table II for females (100 percent above desired level)

A. History of pain and limitation of motion in any weight bearing joint or spine (on physical examination) associated with x-ray evidence of arthritis in a weight bearing joint or spine; or

B. Hypertension with diastolic blood pressure persistently in excess of 100 mm. Hg measured with appropriate size cuff; or

C. History of congestive heart failure manifested by past evidence of vascular congestion such as hepatomegaly, peripheral or pulmonary edema; or

D. Chronic venous insufficiency with superficial varicosities in a lower extremity with pain on weight bearing and persistent edema; or

E. Respiratory disease with total forced vital capacity equal to or less than 2.0 L. or a level of hypoxemia at rest equal to or less than the values specified in Table III-A or III-B or III-C.

### TABLE I—MEN

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### TABLE III—A

[Applicable at test sites less than 3,000 feet above sea level]

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### TABLE III—B

[Applicable at test sites 3,000 through 6,000 feet above sea level]

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### TABLE III—C

[Applicable at test sites over 6,000 feet above sea level]

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### 11.00 NEUROLOGICAL

A. Convulsive disorders. In convulsive disorders, regardless of etiology degree of impairment will be determined according to type, frequency, duration, and sequelae of seizures. At least one detailed description of a typical seizure is required. Such
description includes the presence or absence of aura, tongue bites, sphencter control, injuries associated with the attack, and postictal phenomena. The reporting physician should indicate in the form of paresis or paralysis, tremor, or other involuntary movements, ataxia and sensory disturbances (any or all of which may be due to cerebral cerebellar, brain stem, spinal cord, or peripheral nerve dysfunction) which occur singly or in various combination, frequently provides the sole or partial basis for decision in cases of neurological impairment. The assessment of impairment depends on the degree of interference with locomotion and/or interference with the use of fingers, hands, and arms.

D. In conditions which are episodic in character, such as multiple sclerosis or myasthenia gravis, consideration should be given to frequency and duration of exacerbations, length of remissions, and permanent residua.

E. Multiple sclerosis. The major criteria for evaluating impairment caused by multiple sclerosis are discussed in listing 11.09. Paragraph A provides criteria for evaluating disorganization of motor function and gives reference to 11.04B (11.04B then refers to 11.00C). Paragraph B provides references to other listings for evaluating visual or mental impairments caused by multiple sclerosis. Paragraph C provides criteria for evaluating the impairment of individuals who do not have muscle weakness or other significant disorganization of motor function at rest, but who do develop muscle weakness on activity as a result of fatigue.

Use of the criteria in 11.09C is dependent upon (1) documenting a diagnosis of multiple sclerosis, (2) obtaining a description of fatigue considered to be characteristic of multiple sclerosis, and (3) obtaining evidence that the system has actually become fatigued. The evaluation of the magnitude of the impairment must consider the degree of exercise and the severity of the resulting muscle weakness.

The criteria in 11.09C deals with motor abnormalities which occur on activity. If the disorganization of motor function is present at rest, paragraph A must be used, taking into account any further increase in muscle weakness resulting from activity.

Sensory abnormalities may occur, particularly involving central visual acuity. The decrease in visual acuity may occur after brief attempts at activity involving near vision, such as reading. This decrease in visual acuity may not persist when the specific activity is terminated, as with rest, but is predictably reproduced with resumption of the activity. The impairment of central visual acuity in these cases should be evaluated under the criteria in listing 2.02, taking into account the fact that the decrease in visual acuity will wax and wane.

Clariﬁcation of the evidence regarding central nervous system dysfunction responsible for the symptoms may require supporting technical evidence of functional impairment such as evoked response tests during exercise.
11.01 Category of Impairments, Neurological

11.02 Epilepsy—major motor seizures, (grand mal or psychomotor), documented by EEG and by detailed description of a typical seizure pattern, including all associated phenomena; occurring more frequently than once a month, in spite of at least 3 months of prescribed treatment. With:

A. Daytime episodes (loss of consciousness and convulsive seizures) or

B. Nocturnal episodes manifesting residuals which interfere significantly with activity during the day.

11.03 Epilepsy—minor motor seizures (petit mal, psychomotor, or focal), documented by EEG and by detailed description of a typical seizure pattern, including all associated phenomena; occurring more frequently than once weekly in spite of at least 3 months of prescribed treatment. With alteration of awareness or loss of consciousness and transient postictal manifestations of unconventional behavior or significant interference with activity during the day.

11.04 Central nervous system vascular accident. With one of the following more than 3 months post-vascular accident:

A. Sensory or motor aphasia resulting in ineffective speech or communication; or

B. Significant and persistent disorganization of motor function in two extremities, resulting in sustained disturbance of gross and dexterous movements, or gait and station (see 11.00C).

11.05 Brain tumors.

A. Malignant gliomas (astrocytoma—grades III and IV, glioblastoma multiforme), medulloblastoma, ependymoblastoma, or primary sarcoma; or

B. Astrocytoma (grades I and II), meningioma, pituitary tumors, oligodendrogioma, ependymoma, clivus chordoma, and benign tumors. Evaluate under 11.02, 11.03, 11.04 A, or B, or 12.02.

11.06 Parkinsonian syndrome with the following signs: Significant rigidity, bradykinesia, or tremor in two extremities, which, singly or in combination, result in sustained disturbance of gross and dexterous movements, or gait and station.

11.07 Cerebral palsy. With:

A. IQ of 69 or less; or

B. Abnormal behavior patterns, such as destructiveness or emotional instability; or

C. Significant interference in communication due to speech, hearing, or visual defect; or

D. Disorganization of motor function as described in 11.04B.

11.08 Spinal cord or nerve root lesions, due to any cause with disorganization of motor function as described in 11.04B.

11.09 Multiple sclerosis. With:

A. Disorganization of motor function as described in 11.04B; or

B. Visual or mental impairment as described under the criteria in 2.02, 2.03, 2.04, or 12.02; or

C. Significant, reproducible fatigue of motor function with substantial muscle weakness on repetitive activity, demonstrated on physical examination, resulting from neurological dysfunction in areas of the central nervous system known to be pathologically involved by the multiple sclerosis process.

11.10 Amyotrophic lateral sclerosis. With:

A. Significant bulbar signs; or

B. Disorganization of motor function as described in 11.04B.

11.11 Poliomyelitis. With:

A. Persistent difficulty with swallowing or breathing; or

B. Unintelligible speech; or

C. Disorganization of motor function as described in 11.04B.

11.12 Myasthenia gravis. With:

A. Significant difficulty with speaking, swallowing, or breathing while on prescribed therapy; or

B. Significant motor weakness of muscles of extremities on repetitive activity against resistance while on prescribed therapy.

11.13 Muscular dystrophy with disorganization of motor function as described in 11.04B.


11.15 Tabes dorsalis. With:

A. Tabetic crises occurring more frequently than once monthly; or

B. Unsteady, broad-based or ataxic gait causing significant restriction of mobility substantiated by appropriate posterior column signs.

11.16 Subacute combined cord degeneration (pernicious anemia) with disorganization of motor function as described in 11.04B or 11.15B, not significantly improved by prescribed treatment.

11.17 Degenerative disease not elsewhere such as Huntington's chorea, Friedreich's ataxia, and spino-cerebellar degeneration. With:

A. Disorganization of motor function as described in 11.04B or 11.15B; or

B. Chronic brain syndrome. Evaluate under 12.02.

11.18 Cerebral trauma:

Evaluate under the provisions of 11.02, 11.03, 11.04 and 12.02, as applicable.

11.19 Syringomyelia.

With:

A. Significant bulbar signs; or

B. Disorganization of motor function as described in 11.04B.
The mental disorders listings in 12.00 of the Listing of Impairments will no longer be effective on August 28, 1991, unless extended by the Board or revised and promulgated again.

A. Introduction: The evaluation of disability on the basis of mental disorders requires the documentation of a medically determinable impairment(s) as well as consideration of the degree of limitation such impairment(s) may impose on the individual's ability to work and whether these limitations have lasted or are expected to last for a continuous period of at least 12 months.

The listings for mental disorders are arranged in eight diagnostic categories: organic mental disorders (12.02); schizophrenic, paranoid and other psychotic disorders (12.03); affective disorders (12.04); mental retardation and autism (12.05); anxiety related disorders (12.06); somatoform disorders (12.07); personality disorders (12.08); and substance addiction disorders (12.09).

Each diagnostic group, except listings 12.05 and 12.09, consists of a set of clinical findings (paragraph A criteria), one or more of which must be met, and which, if met, lead to a test of functional restrictions (paragraph B criteria), two or three of which must also be met. There are additional considerations (paragraph C criteria) in listings 12.03 and 12.06, discussed therein.

The purpose of including the criteria in paragraphs B and C of the listings for mental disorders is to medically substantiate the presence of a mental disorder. Specific signs and symptoms under any of the listings 12.02 through 12.09 cannot be considered in isolation from the description of the mental disorder contained at the beginning of each listing category. Impairments should be analyzed or reviewed under the mental category(ies) which is supported by the individual's clinical findings.

The purpose of including the criteria in paragraph A of the listings for mental disorders is to describe those functional limitations associated with mental disorders which are incompatible with the ability to work. The restrictions listed in paragraphs B and C must be the result of the mental disorder which is manifested by the clinical findings outlined in paragraph A. The criteria included in paragraphs B and C of the listings for mental disorders have been chosen because they represent functional areas deemed essential to work. An individual who is severely limited in these areas as the result of an impairment identified in paragraph A is presumed to be unable to work.

The structure of the listing for substance addiction disorders, listing 12.09, is different from that for the other mental disorder listings. Listing 12.09 is structured as a reference listing, that is, it will only serve to indicate which of the other listed mental or physical impairments must be used to evaluate the behavioral or physical changes resulting from regular use of addictive substances.

The listings for mental disorders are so constructed that an individual meeting or equaling the criteria could not reasonably be expected to engage in gainful work activity.

Individuals who have an impairment with a level of severity which does not meet the criteria of the listings for mental disorders may or may not have the residual functional capacity (RFC) which would enable them to engage in substantial gainful work activity. The determination of mental RFC is crucial to the evaluation of an individual's capacity to engage in substantial gainful work activity when the criteria of the listings for mental disorders are not met or equaled but the impairment is nevertheless severe.

RFC may be defined as a multidimensional description of the work-related abilities which an individual retains in spite of medical impairments. RFC complements the criteria in paragraphs B and C of the listings for mental disorders by requiring consideration of an expanded list of work-related capacities which may be impaired by mental disorder when the impairment is severe but does not meet or equal a listed mental disorder. (While RFC may be applicable in most claims, the law specifies that it does not apply to the following special claims categories: disabled title XVI children below age 18, widows, widowers and surviving divorced wives. The impairment(s) of these categories must meet or equal a listed impairment for the individual to be eligible for benefits based on disability.)

B. Need for Medical Evidence: The existence of a medically determinable impairment of the required duration must be established by medical evidence consisting of clinical signs, symptoms and/or laboratory or psychological test findings. These findings may be intermittent or persistent depending on the nature of the disorder. Clinical signs are medically demonstrable phenomena which reflect specific abnormalities of behavior, affect, thought, memory, orientation, or contact with reality. These signs are typically assessed by a psychiatrist or psychologist and/or documented by psychological tests. Symptoms are complaints presented by the individual. Signs and symptoms generally cluster together to constitute recognizable clinical syndromes (mental disorders). Both symptoms and signs which are part of any diagnosed mental disorder must be considered in evaluating severity.

C. Assessment of Severity: For mental disorders, severity is assessed in terms of the functional limitations imposed by the impairment. Functional limitations are assessed using the criteria in paragraph B of
the listings for mental disorders (descriptions of restrictions of activities of daily living; social functioning; concentration, persistence, or pace; and ability to tolerate increased mental demands associated with work or competition). Where “marked” is used as a standard for measuring the degree of limitation, it means more than moderate, but less than extreme. A marked limitation may arise when several activities or functions are impaired or even when only one is impaired, so long as the degree of limitation is such as to seriously interfere with the ability to function independently, appropriately and effectively. Four areas are considered.

1. Activities of daily living include adaptive activities such as cleaning, shopping, cooking, taking public transportation, paying bills, maintaining a residence, caring appropriately for one’s grooming and hygiene, using telephones and directories, using a post office, etc. In the context of the individual’s overall situation, the quality of these activities is judged by their independence, appropriateness and effectiveness. It is necessary to define the extent to which the individual is capable of initiating and participating in activities independent of supervision or direction.

“Marked” is not the number of activities which are restricted but the overall degree of restriction or combination of restrictions which must be judged. For example, a person who is able to cook and clean might still have marked restrictions of daily activities if the person were too fearful to leave the immediate environment of home and neighborhood, hampering the person’s ability to obtain treatment or to travel away from the immediate living environment.

2. Social functioning refers to an individual’s capacity to interact appropriately and communicate effectively with other individuals. Social functioning includes the ability to get along with others, e.g., family members, friends, neighbors, grocery clerks, landlords, bus drivers, etc. Impaired social functioning may be demonstrated by a history of altercations, evictions, firings, fear of strangers, avoidance of interpersonal relationships, social isolation, etc. Strength in social functioning may be demonstrated by an individual’s ability to initiate social contacts with others, communicate clearly with others, interact and actively participate in group activities, etc. Cooperative behaviors, consideration for others, awareness of others’ feelings, and social maturity also need to be considered. Social functioning in work situations may involve interactions with the public, responding appropriately to persons in authority, e.g., supervisors, or cooperative behaviors involving coworkers.

“Marked” is not the number of areas in which social functioning is impaired, but the overall degree of interference in a particular area or combination of areas of functioning. For example, a person who is highly antagonistic, uncooperative or hostile but is tolerated by local storekeepers may nevertheless have marked restrictions in social functioning because that behavior is not acceptable in other social contexts.

3. Concentration, persistence and pace refer to the ability to sustain focused attention sufficiently long to permit the timely completion of tasks commonly found in work settings. In activities of daily living, concentration may be reflected in terms of ability to complete tasks in everyday household routines. Deficiencies in concentration, persistence and pace are best observed in work and work-like settings. Major impairment in this area can often be assessed through direct psychiatric examination and/or psychological testing, although mental status examination or psychological test data alone should not be used to accurately describe concentration and sustained ability to adequately perform work-like tasks. On mental status examinations, concentration is assessed by tasks such as having the individual subtract serial sevens from 100. In psychological tests of intelligence or memory, concentration is assessed through tasks requiring short-term memory or through tasks that must be completed within established time limits. In work evaluations, concentration, persistence, and pace are assessed through such tasks as filing index cards, locating telephone numbers, or disassembling and reassembling objects. Strengths and weaknesses in areas of concentration can be discussed in terms of frequency of errors, time it takes to complete the task, and extent to which assistance is required to complete the task.

4. Deterioration or decompensation in work or work-like settings refers to repeated failure to adapt to stressful circumstances which cause the individual either to withdraw from that situation or to experience exacerbation of signs and symptoms (i.e., decompensation) with an accompanying difficulty in maintaining activities of daily living, social relationships, and/or maintaining concentration, persistence, or pace (i.e., deterioration which may include deterioration of adaptive behaviors). Stresses common to the work environment include decisions, attendance, schedules, completing tasks, interactions with supervisors, interactions with peers, etc.

D. Documentation: The presence of a mental disorder should be documented primarily on the basis of reports from individual providers, such as psychiatrists and psychologists, and facilities such as hospitals and clinics. Adequate descriptions of functional limitations must be obtained from these or other sources which may include programs and facilities where the individual has been observed over a considerable period of time.
Information from both medical and non-medical sources may be used to obtain detailed descriptions of the individual's activities of daily living; social functioning; concentration, speed, and pace; performance, or ability to tolerate increased mental demands. This information can be provided by programs such as community mental health centers, day care centers, sheltered workshops, etc. It can also be provided by others, including family members, who have knowledge of the individual's functioning. In some cases descriptions of activities of daily living or social functioning given by individuals or treating sources may be insufficiently detailed and/or may be in conflict with the clinical picture otherwise observed or described in the examinations or reports. It is necessary to resolve any inconsistencies or gaps that may exist in order to obtain a proper understanding of the individual's functional restrictions.

An individual's level of functioning may vary considerably over time. The level of functioning at a specific time may seem relatively adequate or, conversely, rather poor. Proper evaluation of the impairment must take any variations in level of functioning into account in arriving at a determination of impairment severity over time. Thus, it is vital to obtain evidence from relevant sources over a sufficiently long period prior to the date of adjudication in order to establish the individual's impairment severity. This evidence should include treatment notes, hospital discharge summaries, and work evaluation or rehabilitation progress notes if these are available.

Some individuals may have attempted to work or may actually have worked during the period of time pertinent to the determination of disability. This may have been an independent attempt at work, or it may have been in conjunction with a community mental health or other sheltered program which may have been of either short or long duration. Information concerning the individual's behavior during any attempt to work and the circumstances surrounding termination of the work effort are particularly useful in determining the individual's ability or inability to function in a work setting.

The results of well-standardized psychological tests such as the Wechsler Adult Intelligence Scale (WAIS), the Minnesota Multiphasic Personality Inventory (MMPI), the Rorschach, and the Thematic Apperception Test (TAT), may be useful in establishing the existence of a mental disorder. For example, the WAIS is useful in establishing mental retardation, and the MMPI, Rorschach, and TAT may provide data supporting several other diagnoses. Broad-based neuropsychological assessments using, for example, the Halstead-Reitan or the Luria-Nebraska batteries may be useful in determining brain function deficiencies, particularly in cases involving subtle findings such as may be seen in traumatic brain injury. In addition, the process of taking a standardized test requires concentration, persistence, and the ability to tolerate increased mental demand (stress). This information can be supplied by many of the tests described below.

In cases involving impaired intellectual functioning, a standardized intelligence test, e.g., the WAIS, should be administered and interpreted by a psychologist or psychiatrist qualified by training and experience to perform such an evaluation. In special circumstances, nonverbal measures, such as the Raven Progressive Matrices, the Leiter international scale, or the Arthur adaptation of the Leiter may be substituted.

Identical IQ scores obtained from different tests do not always reflect a similar degree of intellectual functioning. In this connection, it must be noted that on the WAIS, for example, IQs of 79 and below are characteristic of approximately the lowest 2 percent of the general population. In instances where other tests are administered, it would be necessary to convert the IQ to the corresponding percentile rank in the general population in order to determine the actual degree of impairment reflected by those IQ scores.

In cases where more than one IQ is customarily derived from the test administered, i.e., where verbal, performance, and full-scale IQs are provided as on the WAIS, the lowest of these is used in conjunction with listing 12.05.

In cases where the nature of the individual's intellectual impairment is such that standard intelligence tests, as described above, are precluded, medical reports specifically describing the level of intellectual, social, and physical function should be obtained. Actual observations by Social Security Administration or State agency personnel, reports from educational institutions and information furnished by public welfare agencies or other reliable objective sources should be considered as additional evidence.

E. Chronic Mental Impairments: Particular problems are often involved in evaluating mental impairments in individuals who have long histories of repeated hospitalizations or prolonged outpatient care with supportive therapy and medication. Individuals with chronic psychotic disorders commonly have their lives structured in such a way as to minimize stress and reduce their signs and symptoms. Such individuals may be much more impaired for work than their signs and symptoms would indicate. The results of a
single examination may not adequately describe these individuals' sustained ability to function. It is, therefore, vital to review all pertinent information relative to the individual's condition, especially at times of increased stress. It is mandatory to attempt to obtain adequate descriptive information from all sources which have treated the individual or in the time period relevant to the decision.

F. Effects of Structured Settings: Particularly in cases involving chronic mental disorders, overt symptomatology may be controlled or attenuated by psychosocial factors such as placement in a hospital, board and care facility, or other environment that provides similar structure. Highly structured and supportive settings may greatly reduce the mental demands placed on an individual. With lowered mental demands, overt signs and symptoms of the underlying mental disorder may be minimized. At the same time, however, the individual's ability to function outside of such a structured and/or supportive setting may not have changed. An evaluation of individuals whose symptomatology is controlled or attenuated by psychosocial factors must consider the ability of the individual to function outside of such highly structured settings. (For these reasons the paragraph C criteria were added to Listings 12.03 and 12.06.)

G. Effects of Medication: Attention must be given to the effect of medication on the individual's signs, symptoms and ability to function. While psychotropic medications may control certain primary manifestations of a mental disorder, e.g., hallucinations, such treatment may or may not affect the functional limitations imposed by the mental disorder. In cases where overt symptomatology is attenuated by the psychotropic medications, particular attention must be focused on the functional restrictions which may persist. These functional restrictions are also to be used as the measure of impairment severity. (See the paragraph C criteria in Listings 12.03 and 12.06.)

Neuroleptics, the medicines used in the treatment of some mental illnesses, may cause dryness, blunted affect, or other side effects involving other body systems. Such side effects must be considered in evaluating overall impairment severity. Where adverse effects of medications contribute to the impairment severity and the impairment does not meet or equal the listings but is nonetheless severe, such adverse effects must be considered in the assessment of the mental residual functional capacity.

H. Effect of Treatment: It must be remembered that with adequate treatment some individuals suffering with chronic mental disorders not only have their symptoms and signs ameliorated but also return to a level of function close to that of their premorbid status. Our discussion here in 12.00H has been designed to reflect the fact that present day treatment of a mentally impaired individual may or may not assist in the achievement of an adequate level of adaptation required in the work place. (See the paragraph C criteria in Listings 12.03 and 12.06.)

1. Technique for Reviewing the Evidence in Mental Disorders Claims to Determine Level of Impairment Severity: A special technique has been developed to ensure that all evidence needed for the evaluation of impairment severity in claims involving mental impairment is obtained, considered and properly evaluated. This technique, which is used in connection with the sequential evaluation process, is explained in §404.1520a and §416.920a.

12.01 Category of Impairments-Mental
12.02 Organic Mental Disorders: Psychological or behavioral abnormalities associated with a dysfunction of the brain. History and physical examination or laboratory tests demonstrate the presence of a specific organic factor judged to be etiologically related to the abnormal mental state and loss of previously acquired functional abilities. The required level of severity for these disorders is met when the requirements in both A and B are satisfied.

A. Demonstration of a loss of specific cognitive abilities or affective changes and the medically documented persistence of at least one of the following:
1. Disorientation to time and place; or
2. Memory impairment, either short-term (inability to learn new information), intermediate, or long-term (inability to remember information that was known sometime in the past); or
3. Perceptual or thinking disturbances (e.g., hallucinations, delusions); or
4. Change in personality; or
5. Disturbance in mood; or
6. Emotional lability (e.g., explosive temper outbursts, sudden crying, etc.) and impairment in impulse control; or
7. Loss of measured intellectual ability of at least 15 I.Q. points from premorbid levels or overall impairment index clearly within the severely impaired range on neuropsychological testing, e.g., the Luria-Nebraska, Halstead-Reitan, etc.; AND

B. Resulting in at least two of the following:
1. Marked restriction of activities of daily living; or
2. Marked difficulties in maintaining social functioning; or
3. Deficiencies of concentration, persistence or pace resulting in frequent failure to complete tasks in a timely manner (in work settings or elsewhere); or
4. Repeated episodes of deterioration or decompensation in work or work-like settings which cause the individual to withdraw from that situation or to experience exacerbation
Psychotic Disorders:

Affective Disorders:

Mental Retardation and Autism:

Railroad Retirement Board

of signs and symptoms (which may include deterioration of adaptive behaviors).

Psychotic Disorders: Characterized by the onset of psychotic features with deterioration from a previous level of functioning.

The required level of severity for these disorders is met when the requirements in both A and B are satisfied, or when the requirements in C are satisfied.

A. Medically documented persistence, either continuous or intermittent, of one or more of the following:
   1. Delusions or hallucinations; or
   2. Catatonic or other grossly disorganized behavior; or
   3. Incoherence, loosening of associations, illogical thinking, or poverty of content of speech if associated with one of the following:
      a. Blunt affect; or
      b. Flat affect; or
      c. Inappropriate affect; or
   4. Emotional withdrawal and/or isolation;

   AND

   B. Resulting in at least two of the following:
      1. Marked restriction of activities of daily living; or
      2. Marked difficulties in maintaining social functioning; or
      3. Deficiencies of concentration, persistence or pace resulting in frequent failure to complete tasks in a timely manner (in work settings or elsewhere); or
      4. Repeated episodes of deterioration or decompensation in work or work-like settings which cause the individual to withdraw from that situation or to experience exacerbation of signs and symptoms (which may include deterioration of adaptive behaviors);

   OR

C. Medically documented history of one or more episodes of acute symptoms, signs and functional limitations which at the time met the requirements in A and B of this listing, although these symptoms or signs are currently attenuated by medication or psychosocial support, and one of the following:

1. Repeated episodes of deterioration or decompensation in situations which cause the individual to withdraw from that situation or to experience exacerbation of signs or symptoms (which may include deterioration of adaptive behaviors); or

2. Documented current history of two or more years of inability to function outside of a highly supportive living situation.

12.04 Affective Disorders: Characterized by a disturbance of mood, accompanied by a full or partial manic or depressive syndrome. Mood refers to a prolonged emotion that colors the whole psychic life; it generally involves either depression or elation.

The required level of severity for these disorders is met when the requirements in both A and B are satisfied.

A. Medically documented persistence, either continuous or intermittent, of one of the following:
   1. Depressive syndrome characterized by at least four of the following:
      a. Anhedonia or pervasive loss of interest in almost all activities; or
      b. Appetite disturbance with change in weight; or
      c. Sleep disturbance; or
      d. Psychomotor agitation or retardation; or
      e. Decreased energy; or
      f. Feelings of guilt or worthlessness; or
      g. Difficulty concentrating or thinking; or
      h. Thoughts of suicide; or
      i. Hallucinations, delusions or paranoid thinking; or
   2. Manic syndrome characterized by at least three of the following:
      a. Hyperactivity; or
      b. Pressure of speech; or
      c. Flight of ideas; or
      d. Inflated self-esteem; or
      e. Decreased need for sleep; or
      f. Easy distractibility; or
      g. Involvement in activities that have a high probability of painful consequences which are not recognized; or
      h. Hallucinations, delusions or paranoid thinking; or
   3. Bipolar syndrome with a history of episodic periods manifested by the full symptomatic picture of both manic and depressive syndromes (and currently characterized by either or both syndromes);

   AND

   B. Resulting in at least two of the following:
      1. Marked restriction of activities of daily living; or
      2. Marked difficulties in maintaining social functioning; or
      3. Deficiencies of concentration, persistence or pace resulting in frequent failure to complete tasks in a timely manner (in work settings or elsewhere); or
      4. Repeated episodes of deterioration or decompensation in work or work-like settings which cause the individual to withdraw from that situation or to experience exacerbation of signs or symptoms (which may include deterioration of adaptive behaviors).

12.05 Mental Retardation and Autism: Mental retardation refers to a significantly subaverage general intellectual functioning with deficits in adaptive behavior initially manifested during the developmental period (before age 22). (Note: The scores specified below refer to those obtained on the WAIS, and are used only for reference purposes. Scores obtained on other standardized and
Somatoform Disorders: Physical symptoms for which there are no demonstrable organic findings or known physiological mechanisms.

The required level of severity for these disorders is met when the requirements in both A and B are satisfied.

A. Medically documented evidence of one of the following:
1. A history of multiple physical symptoms of several years duration, beginning before age 30, that have caused the individual to take medicine frequently, see a physician often and alter life patterns significantly; or
2. Persistent nonorganic disturbance of one of the following:
   a. Vision; or
   b. Speech; or
   c. Hearing; or
   d. Use of a limb; or
   e. Movement and its control (e.g., coordination disturbance, psychogenic seizures, akinesia, dyskinesia; or
   f. Sensation (e.g., diminished or heightened).

3. Unrealistic interpretation of physical signs or sensations associated with the preoccupation or belief that one has a serious disease or injury; AND

Anxiety Related Disorders: In these disorders anxiety is either the predominant disturbance or it is experienced if the individual attempts to master symptoms; for example, confronting the dreaded object or situation in a phobic disorder or resisting the obsessions or compulsions in obsessive compulsive disorders.

The required level of severity for these disorders is met when the requirements in both A and B are satisfied, or when the requirements in both A and C are satisfied.

A. Mental incapacity evidenced by dependence upon others for personal needs (e.g., toileting, eating, dressing, or bathing) and inability to follow directions, such that the use of standardized measures of intellectual functioning is precluded;
OR
B. A valid verbal, performance, or full scale IQ of 59 or less;
OR
C. A valid verbal, performance, or full scale IQ of 60 to 69 inclusive or in the case of autism gross deficits of social and communicative skills with two of the following:
   1. Marked restriction of activities of daily living; or
   2. Marked difficulties in maintaining social functioning; or
   3. Deficiencies of concentration, persistence or pace resulting in frequent failure to complete tasks in a timely manner (in work settings or elsewhere); or
4. Repeated episodes of deterioration or decompensation in work or work-like settings which cause the individual to withdraw from that situation or to experience exacerbation of signs and symptoms (which may include deterioration of adaptive behaviors);
OR
C. Resulting in complete inability to function independently outside the area of one's home.

12.06 Anxiety Related Disorders: In these disorders anxiety is either the predominant disturbance or it is experienced if the individual attempts to master symptoms; for example, confronting the dreaded object or situation in a phobic disorder or resisting the obsessions or compulsions in obsessive compulsive disorders.

The required level of severity for these disorders is met when the requirements in both A and B are satisfied.

A. Medically documented findings of at least one of the following:
   1. Generalized persistent anxiety accompanied by three out of four of the following signs or symptoms:
      a. Motor tension; or
      b. Autonomic hyperactivity; or
      c. Apprehensive expectation; or
      d. Vigilance and scanning;

   2. A persistent irrational fear of a specific object, activity, or situation which results in a compelling desire to avoid the dreaded object, activity, or situation; or
   3. Recurrent severe panic attacks manifested by a sudden unpredictable onset of intense apprehension, fear, terror and sense of impending doom occurring on the average of at least once a week; or
   4. Recurrent obsessions or compulsions which are a source of marked distress; or
   5. Recurrent and intrusive recollections of a traumatic experience, which are a source of marked distress; AND

B. Resulting in at least two of the following:
   1. Marked restriction of activities of daily living; or
   2. Marked difficulties in maintaining social functioning; or
   3. Deficiencies of concentration, persistence or pace resulting in frequent failure to complete tasks in a timely manner (in work settings or elsewhere); or
4. Repeated episodes of deterioration or decompensation in work or work-like settings which cause the individual to withdraw from that situation or to experience exacerbation of signs and symptoms (which may include deterioration of adaptive behaviors);
OR
C. Resulting in complete inability to function independently outside the area of one's home.

12.07 Somatoform Disorders: Physical symptoms for which there are no demonstrable organic findings or known physiological mechanisms.

The required level of severity for these disorders is met when the requirements in both A and B are satisfied.

A. Medically documented evidence of one of the following:
1. A history of multiple physical symptoms of several years duration, beginning before age 30, that have caused the individual to take medicine frequently, see a physician often and alter life patterns significantly; or
2. Persistent nonorganic disturbance of one of the following:
   a. Vision; or
   b. Speech; or
   c. Hearing; or
   d. Use of a limb; or
   e. Movement and its control (e.g., coordination disturbance, psychogenic seizures, akinesia, dyskinesia; or
   f. Sensation (e.g., diminished or heightened).

3. Unrealistic interpretation of physical signs or sensations associated with the preoccupation or belief that one has a serious disease or injury; AND
B. Resulting in three of the following:
1. Marked restriction of activities of daily living; or
2. Marked difficulties in maintaining social functioning; or
3. Deficiencies of concentration, persistence or pace resulting in frequent failure to complete tasks in a timely manner (in work settings or elsewhere); or
4. Repeated episodes of deterioration or decompensation in work or work-like settings which cause the individual to withdraw from that situation or to experience exacerbation of signs and symptoms (which may include deterioration of adaptive behavior).

12.08 Personality Disorders: A personality disorder exists when personality traits are inflexible and maladaptive and cause either significant impairment in social or occupational functioning or subjective distress. Characteristic features are typical of the individual's long-term functioning and are not limited to discrete episodes of illness. The required level of severity for these disorders is met when the requirements in both A and B are satisfied.

A. Deeply ingrained, maladaptive patterns of behavior associated with one of the following:
1. Seclusiveness or autistic thinking; or
2. Pathologically inappropriate suspiciousness or hostility; or
3. Oddities of thought, perception, speech and behavior; or
4. Persistent disturbances of mood or affect; or
5. Pathological dependence, passivity, or aggressivity; or
6. Intense and unstable interpersonal relationships and impulsive and damaging behavior.

AND

B. Resulting in three of the following:
1. Marked restriction of activities of daily living; or
2. Marked difficulties in maintaining social functioning; or
3. Deficiencies of concentration, persistence or pace resulting in frequent failure to complete tasks in a timely manner (in work settings or elsewhere); or
4. Repeated episodes of deterioration or decompensation in work or work-like settings which cause the individual to withdraw from that situation or to experience exacerbation of signs and symptoms (which may include deterioration of adaptive behavior).

12.09 Substance Addiction Disorders: Behavioral changes or physical changes associated with the regular use of substances that affect the central nervous system. The required level of severity for these disorders is met when the requirements in any of the following (A through I) are satisfied.

A. Organic mental disorders. Evaluate under 12.02.
B. Depressive syndrome. Evaluate under 12.04.
C. Anxiety disorders. Evaluate under 12.06.
D. Personality disorders. Evaluate under 12.08.
F. Liver damage. Evaluate under 5.05.
G. Gastritis. Evaluate under 5.04.
H. Pancreatitis. Evaluate under 5.08.
I. Seizures. Evaluate under 11.02 or 11.03.

13.00 NEOPLASTIC DISEASES, MALIGNANT

A. Introduction: The determination of the level of impairment resulting from malignant tumors is made from a consideration of the site of the lesion, the histogenesis of the tumor, the extent of involvement, the apparent adequacy and response to therapy (surgery, irradiation, hormones, chemotherapy, etc.), and the magnitude of the post therapeutic residuals.

B. Documentation: The diagnosis of malignant tumors should be established on the basis of symptoms, signs, and laboratory findings. The site of the primary, recurrent, and metastatic lesion must be specified in all cases of malignant neoplastic diseases. If an operative procedure has been performed, the evidence should include a copy of the operative note and the report of the gross and microscopic examination of the surgical specimen. If these documents are not obtainable, then the summary of hospitalization or report from the treating physician must include details of the findings at surgery and the results of the pathologist's gross and microscopic examination of the tissues. For those cases in which a disabling impairment was not established when therapy was begun but progression of the disease is likely, current medical evidence should include a report of a recent examination directed especially at local or regional recurrence, soft part or skeletal metastases, and significant posttherapeutic residuals.

C. Evaluation. Usually, when the malignant tumor consists of a local lesion with metastases to the regional lymph nodes which apparently has been completely excised, imminent recurrence or metastases is not anticipated. A number of exceptions are noted in the specific Listings. For adjudicative purposes, “distant metastases” or “metastases beyond the regional lymph nodes” refers to metastasis beyond the lines of the usual radical en bloc resection. Local or regional recurrence after radical surgery or pathological evidence of incomplete excision by radical surgery is to be equated with unresectable lesions (except for carcinoma of the breast, 13.09c) and, for the purposes of our program, may be evaluated as “inoperable.”
Local or regional recurrence after incomplete excision of a localized and still completely resectable tumor is not to be equated with recurrence after radical surgery. In the evaluation of lymphomas, the tissue type and site of involvement are not necessarily indicators of the degree of impairment.

When a malignant tumor has metastasized beyond the regional lymph nodes, the impairment will usually be found to meet the requirements of a specific listing. Exceptions are hormone-dependent tumors, isotope-sensitive metastases, and metastases from seminoma of the testicles which are controlled by definitive therapy.

When the original tumor and any metastases have apparently disappeared and have not been evident for 3 or more years, the impairment does not meet the criteria under this body system.

D. Effects of therapy. Significant posttherapeutic residuals, not specifically included in the category of impairments for malignant neoplasms, should be evaluated according to the affected body system.

Where the impairment is not listed in the Listing of Impairments and is not medically equivalent to a listed impairment, the impact of any residual impairment including that caused by therapy must be considered.

The therapeutic regimen and consequent adverse response to therapy may vary widely; therefore, each case must be considered on an individual basis. It is essential to obtain a specific description of the therapeutic regimen, including the drugs given, dosage, frequency of drug administration, and plans for continued drug administration. It is necessary to obtain a description of the complications or any other adverse response to therapy such as nausea, vomiting, diarrhea, weakness, dermatologic disorders, or reactive mental disorders. Since the severity of the adverse effects of anticancer chemotherapy may change during the period of drug administration, the decision regarding the impact of drug therapy should be based on a sufficient period of therapy to permit proper consideration.

E. Onset. To establish onset of disability prior to the time a malignancy is first demonstrated to be inoperable or beyond control by other modes of therapy (and prior evidence is nonexistent) requires medical judgment based on medically reported symptoms, the type of the specific malignancy, its location, and extent of involvement when first demonstrated.

13.01 Category of Impairments, Neoplastic Diseases—Malignant

13.02 Head and neck (except salivary glands—13.07, thyroid gland—13.08, and mandible, maxilla, orbit, or temporal fossa—13.11):

A. Inoperable; or
B. Not controlled by prescribed therapy; or
C. Recurrent after radical surgery or irradiation; or
D. With distant metastases; or
E. Epidermoid carcinoma occurring in the pyriform sinus or posterior third of the tongue.

13.03 Sarcoma of skin:

A. Angiosarcoma with metastases to regional lymph nodes or beyond; or
B. Mycosis fungoides with metastases to regional lymph nodes, or with visceral involvement.

13.04 Sarcoma of soft parts: Not controlled by prescribed therapy.

13.05 Malignant melanoma:

A. Recurrent after wide excision; or
B. With metastases to adjacent skin (satellite lesions) or elsewhere.

13.06 Lymph nodes:

A. Hodgkin’s disease or non-Hodgkin’s lymphoma with progressive disease not controlled by prescribed therapy; or
B. Metastatic carcinoma in a lymph node (except for epidermoid carcinoma in a lymph node in the neck) where the primary site is not determined after adequate search; or
C. Epidermoid carcinoma in a lymph node in the neck not responding to prescribed therapy.

13.07 Salivary glands—carcinoma or sarcoma with metastases beyond the regional lymph nodes.

13.08 Thyroid gland—carcinoma with metastases beyond the regional lymph nodes, not controlled by prescribed therapy.

13.09 Breast:

A. Inoperable carcinoma; or
B. Inflammatory carcinoma; or
C. Recurrent carcinoma, except local recurrence controlled by prescribed therapy; or
D. Distant metastases from breast carcinoma (bilateral breast carcinoma, synchronous or metachronous is usually primary in each breast); or
E. Sarcoma with metastases anywhere.

13.10 Skeletal system (exclusive of the jaw): not controlled by prescribed therapy.

13.11 Mandible, maxilla, orbit, or temporal fossa:

A. Sarcoma of any type with metastases; or
B. Carcinoma of the antrum with extension into the orbit or ethmoid or sphenoid sinus, or with regional or distant metastases; or
C. Orbital tumors with intracranial extension; or
D. Tumors of the temporal fossa with perforation of skull and meningeal involvement; or
E. Adamantinoma with orbital or intracranial infiltration; or
F. Tumors of Rathke’s pouch with infiltration of the base of the skull or metastases.

13.12 Brain or spinal cord:
A. Metastatic carcinoma to brain or spinal cord.
B. Evaluate other tumors under the criteria described in 11.05 and 11.08.

13.13 Lungs:
A. Unresectable or with incomplete excision; or
B. Recurrence or metastases after resection; or
C. Oat cell (small cell) carcinoma; or
D. Squamous cell carcinoma, with metastases beyond the hilar lymph nodes; or
E. Other histologic types of carcinoma, including undifferentiated and mixed-cell types (but excluding oat cell carcinoma, 13.13C, and squamous cell carcinoma, 13.13D), with metastases to the hilar lymph nodes.

13.14 Pleura or mediastinum:
A. Malignant mesothelioma of pleura; or
B. Malignant tumors, metastatic to pleura; or
C. Malignant primary tumor of the mediastinum not controlled by prescribed therapy.

13.15 Abdomen:
A. Generalized carcinomatosis; or
B. Retropertitoneal cellular sarcoma not controlled by prescribed therapy; or
C. Ascites with demonstrated malignant cells.

13.16 Esophagus or stomach:
A. Carcinoma or sarcoma of the esophagus; or
B. Carcinoma of the stomach with metastases to the regional lymph nodes or extension to surrounding structure; or
C. Sarcoma of stomach not controlled by prescribed therapy; or
D. Inoperable carcinoma; or
E. Recurrence or metastases after resection.

13.17 Small intestine:
A. Carcinoma, sarcoma, or carcinoid tumor with metastases beyond the regional lymph nodes; or
B. Recurrence of carcinoma, sarcoma, or carcinoid tumor after resection; or
C. Sarcoma, not controlled by prescribed therapy.

13.18 Large intestine (from ileocecal valve to and including anal canal)—carcinoma or sarcoma:
A. Unresectable; or
B. Metastases beyond the regional lymph nodes; or
C. Recurrence or metastases after resection.

13.19 Liver or gallbladder:
A. Primary or metastatic malignant tumors of the liver; or
B. Carcinoma of the gallbladder; or
C. Carcinoma of the bile ducts.

13.20 Pancreas:
A. Carcinoma except islet cell carcinoma; or
B. Islet cell carcinoma which is unresectable and physiologically active.

13.21 Kidneys, adrenal glands, or ureters—carcinoma:
A. Unresectable; or
B. With hematogenous spread to distant sites; or
C. With metastases to regional lymph nodes.

13.22 Urinary bladder—carcinoma. With:
A. Infiltration beyond the bladder wall; or
B. Metastases to regional lymph nodes; or
C. Unresectable; or
D. Recurrence after total cystectomy; or
E. Evaluate renal impairment after total cystectomy under the criteria in 6.02.

13.23 Prostate gland—carcinoma not controlled by prescribed therapy.

13.24 Testicles:
A. Choriocarcinoma; or
B. Other malignant primary tumors with progressive disease not controlled by prescribed therapy.

13.25 Uterus—carcinoma or sarcoma (corpus or cervix):
A. Inoperable and not controlled by prescribed therapy; or
B. Metastases to regional lymph nodes; or
C. Unresectable infiltration; or
D. Unresectable metastases to omentum or elsewhere in the peritoneal cavity; or
E. With distant metastases.

13.26 Ovaries—all malignant, primary or recurrent tumors. With:
A. Ascites with demonstrated malignant cells; or
B. Unresectable infiltration; or
C. Unresectable metastases to omentum or elsewhere in the peritoneal cavity; or
D. Distant metastases.

13.27 Leukemia: Evaluate under the criteria of 7.00ff, Hemic and Lymphatic System.

13.28 Uterine (Fallopian) tubes—carcinoma or sarcoma:
A. Unresectable, or
B. Metastases to regional lymph nodes.

13.29 Penis—carcinoma with metastases to regional lymph nodes.

13.30 Vulva—carcinoma, with distant metastases.

Part B
Medical criteria for the evaluation of impairments of children under age 18 (where criteria in Part A do not give appropriate consideration to the particular disease process in childhood).

Sec.
100.00 Growth Impairment.
101.00 Musculoskeletal System.
102.00 Special Senses and Speech.
103.00 Respiratory System.
104.00 Cardiovascular System.
105.00 Digestive System.
106.00 Genito-Urinary System.
107.00 Hemic and Lymphatic System.
108.00 [Reserved]
109.00 Endocrine System.
GROWTH IMPAIRMENT

- Impairment of growth may be disabling in itself or it may be an indicator of the severity of the impairment due to a specific disease process. Determinations of growth impairment should be based on the comparison of current height with at least three previous determinations, including length at birth, if available. Heights (or lengths) should be plotted on a standard growth chart, such as derived from the National Center for Health Statistics: NCHS Growth Charts. Height should be measured without shoes. Body weight corresponding to the ages represented by the heights should be furnished. This will provide a basis upon which to identify those children whose short stature represents a familial characteristic rather than a result of disease. This is particularly true for adjudication under 100.02B.

- Bone age determinations should include a full descriptive report of roentgenograms specifically obtained to determine bone age and must cite the standardization method used. Where roentgenograms must be obtained currently as a basis for adjudication under 100.03, views of the left hand and wrist should be ordered. In addition, roentgenograms of the knee and ankle should be obtained when cessation of growth is being evaluated in an older child at, or past, puberty.

- The criteria in this section are applicable until closure of the major epiphyses. The cessation of significant increase in height at that point would prevent the application of these criteria.

- It is not identified as growth impairment due to deformity or musculoskeletal disease at the birth of the child.

- The measurements of joint motion, not identified as resulting from a specific stage of many skeletal diseases and conditions, such as traumatic arthritis, collagen disorders, septic arthritis, congenital dislocation of the hip, aseptic necrosis of the hip, slipped capital femoral epiphyses, skeletal dysplasias, etc.

100.00 Category of Impairments, Musculoskeletal

101.00 MUSCULOSKELETAL SYSTEM

A. Rheumatoid arthritis. Documentation of the diagnosis of juvenile rheumatoid arthritis should be made according to an established protocol, such as that published by the Arthritis Foundation, Bulletin on the Rheumatic Diseases, Vol. 23, 1972-1973 Series, p 712. Inflammatory signs include persistent pain, tenderness, erythema, swelling, and increased local temperature of a joint. B. The measurements of joint motion are based on the technique for measurements described in the "Joint Method of Measuring and Recording," published by the American Academy of Orthopedic Surgeons in 1965, or "The Extremities and Back" in Guides to the Evaluation of Permanent Impairment, Chicago, American Medical Association, 1971, Chapter 1, pp. 1-48.

C. Degenerative arthritis may be the end stage of many skeletal diseases and conditions, such as traumatic arthritis, collagen disorders, septic arthritis, congenital dislocation of the hip, aseptic necrosis of the hip, slipped capital femoral epiphyses, skeletal dysplasias, etc.

101.01 Category of Impairments, Musculoskeletal

101.02 Juvenile rheumatoid arthritis. With:

- A. Persistence or recurrence of joint inflammation despite three months of medical treatment and one of the following: 1. Limitation of motion of two major joints of 50 percent or greater; or 2. Fixed deformity of two major weight-bearing joints of 30 degrees or more; or 3. Radiographic changes of joint narrowing, erosion, or subluxation; or 4. Persistent or recurrent systemic involvement such as iridocyclitis or pericarditis; or

- B. Steroid dependence.

101.03 Deficit of musculoskeletal function due to deformity or musculoskeletal disease and one of the following:

- A. Walking is markedly reduced in speed or distance despite orthotic or prosthetic devices; or

- B. Ambulation is possible only with obligatory bilateral upper limb assistance (e.g., with walker, crutches); or

- C. Inability to perform age-related personal self-care activities involving feeding, dressing, and personal hygiene.

101.05 Disorders of the spine.

A. Fracture of vertebra with cord involvement (substantiated by appropriate sensory and motor loss); or

B. Scoliosis (congenital idiopathic or neuromyopathic). With:

- A. Major spinal curve measuring 60 degrees or greater; or

- B. Spinal fusion of six or more levels. Consider under a disability for one year from the time of surgery; thereafter evaluate the residual impairment; or

- C. FEV (vital capacity) of 50 percent or less of predicted normal values for the individual's measured (actual) height; or

101.00 MUSCULOSKELETAL SYSTEM

A. Rheumatoid arthritis. Documentation of the diagnosis of juvenile rheumatoid arthritis should be made according to an established protocol, such as that published by the Arthritis Foundation, Bulletin on the Rheumatic Diseases, Vol. 23, 1972-1973 Series, p 712. Inflammatory signs include persistent pain, tenderness, erythema, swelling, and increased local temperature of a joint. B. The measurements of joint motion are based on the technique for measurements described in the "Joint Method of Measuring and Recording," published by the American Academy of Orthopedic Surgeons in 1965, or "The Extremities and Back" in Guides to the Evaluation of Permanent Impairment, Chicago, American Medical Association, 1971, Chapter 1, pp. 1-48.

C. Degenerative arthritis may be the end stage of many skeletal diseases and conditions, such as traumatic arthritis, collagen disorders, septic arthritis, congenital dislocation of the hip, aseptic necrosis of the hip, slipped capital femoral epiphyses, skeletal dysplasias, etc.
Railroad Retirement Board

C. Kyphosis or lordosis measuring 90 degrees or greater.

101.08 Chronic osteomyelitis with persistence or recurrence of inflammatory signs or drainage for at least 6 months despite prescribed therapy and consistent radiographic findings.

102.00 Special Senses and Speech

A. Visual impairments in children. Impairment of central visual acuity should be determined with use of the standard Snellen test chart. Where this cannot be used, as in very young children, a complete description should be provided of the findings using other appropriate methods of examination, including a description of the techniques used for determining the central visual acuity for distance.

The accommodative reflex is generally not present in children under 6 months of age. In premature infants, it may not be present until 6 months plus the number of months the child is premature. Therefore absence of accommodative reflex will be considered as indicating a visual impairment only in children above this age (6 months).

Documentation of a visual disorder must include description of the ocular pathology.

B. Hearing impairments in children. The criteria for hearing impairments in children take into account that a lesser impairment in hearing which occurs at an early age may result in a severe speech and language disorder.

Improvement by a hearing aid, as predicted by the testing procedure, must be demonstrated to be feasible in that child, since younger children may be unable to use a hearing aid effectively.

The type of audiometric testing performed must be described and a copy of the results must be included. The pure tone air conduction hearing levels in 102.08 are based on American National Standard Institute Specifications for Audiometers, S3.6-1969 (ANSI-1969). The report should indicate the specifications used to calibrate the audiometer.

The finding of a severe impairment will be based on the average hearing levels at 500, 1000, 2000, and 3000 Hertz (Hz) in the better ear, and on speech discrimination, as specified in §102.08.

102.01 Category of Impairments, Special Sense Organs

102.02 Impairments of central visual acuity.

A. Remaining vision in the better eye after best correction is 20/200 or less; or

B. For children under 3 years of age at time of adjudication:

1. Absence of accommodative reflex (see 102.00A for exclusion of children under 6 months of age); or

2. Retrolental fibroplasia with macular scarring or neovascularization; or

3. Bilateral congenital cataracts with visualization of retinal red reflex only or when associated with other ocular pathology.

102.08 Hearing impairments.

A. For children below 5 years of age at time of adjudication, inability to hear air conduction thresholds at an average of 40 decibels (db) hearing level or greater in the better ear; or

B. For children 5 years of age and above at time of adjudication:

1. Inability to hear air conduction thresholds at an average of 70 decibels (db) or greater in the better ear; or

2. Speech discrimination scores at 40 percent or less in the better ear; or

3. Inability to hear air conduction thresholds at an average of 40 decibels (db) or greater in the better ear, and a speech and language disorder which significantly affects the clarity and content of the speech and is attributable to the hearing impairment.

103.00 Respiratory System

A. Documentation of pulmonary insufficiency. The reports of spirometric studies for evaluation under Table I must be expressed in liters (BTPS). The reported FEV1 must represent the largest of at least three satisfactory attempts. The appropriately labeled spirometric tracing of three FEV1 maneuvers must be submitted with the report, showing distance per second on the abscissa and distance per liter on the ordinate. The unit distance for volume on the tracing should be at least 20 mm per liter and the paper speed at least 20 mm per second. The height of the individual without shoes must be recorded.

The ventilatory function studies should not be performed during or soon after an acute episode or exacerbation of a respiratory illness. In the presence of acute bronchospasm, or where the FEV1 is less than that stated in Table I, the studies should be repeated after the administration of a nebulized bronchodilator. If a bronchodilator was not used in such instances, the reason should be stated in the report.

A statement should be made as to the child's ability to understand directions and to cooperate in performance of the test, and to cooperate in performance of the test, and should include an evaluation of the child's effort. When tests cannot be performed or completed, the reason (such as a child's young age) should be stated in the report.

B. Cystic fibrosis. This section discusses only the pulmonary manifestations of cystic fibrosis. Other manifestations, complications, or associated disease must be evaluated under the appropriate section.

The diagnosis of cystic fibrosis will be based upon appropriate history, physical examination, and pertinent laboratory findings. Confirmation based upon elevated concentration of sodium or chloride in the sweat should be included, with indication of the technique used for collection and analysis.
A. Recent, recurrent intense asthmatic attacks requiring parenteral medication; or
B. Persistent prolonged expiration with wheezing between acute attacks and radiographic findings of peribronchial disease.

103.13 Pulmonary manifestations of cystic fibrosis. With:
A. FEV₁, equal to or less than the values specified in Table I (see §103.00A for requirements of ventilatory function testing); or
B. For children where ventilatory function testing cannot be performed:
   1. History of dyspnea on mild exertion or chronic frequent productive cough; and
   2. Persistent or recurrent abnormal breath sounds, bilateral rales or rhonchi; and
   3. Radiographic findings of extensive disease with hyperaeration and bilateral peribronchial infiltration.

<table>
<thead>
<tr>
<th>TABLE I</th>
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<tbody>
<tr>
<td><strong>Height (in centimeters)</strong></td>
</tr>
<tr>
<td>110 or less</td>
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<tr>
<td>120</td>
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<tr>
<td>130</td>
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<tr>
<td>140</td>
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<tr>
<td>150</td>
</tr>
<tr>
<td>160</td>
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<tr>
<td>170 or more</td>
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</table>

104.00 Cardiovascular System
A. General. Evaluation should be based upon history, physical findings, and appropriate laboratory data. Reported abnormalities should be consistent with the pathologic diagnosis. The actual electrocardiographic tracing, or an adequate marked photopy, must be included. Reports of other pertinent studies necessary to substantiate the diagnosis or describe the severity of the impairment must also be included:
B. Evaluation of cardiovascular impairment in children requires two steps:
   1. The delineation of a specific cardiovascular disturbance, either congenital or acquired. This may include arterial or venous disease, rhythm disturbance, or disease involving the valves, septa, myocardium or pericardium; and
   2. Documentation of the severity of the impairment, with medically determinable and consistent cardiovascular signs, symptoms, and laboratory data. In cases where impairment characteristics are unquestionably secondary to the cardiovascular disturbance, additional documentation of the severity of the impairment (e.g., catheterization data, if performed) will be necessary.
C. Chest roentgenogram (6 ft. PA film) will be considered indicative of cardiomegaly if:
   1. The cardiothoracic ratio is over 60 percent at age one year or less, or 55 percent at more than one year of age; or
   2. The cardiac size is increased over 15 percent from any prior chest roentgenograms; or
   3. Specific chamber or vessel enlargement is documented in accordance with established criteria.
D. Tables I, II, and III below are designed for case adjudication and not for diagnostic purposes. The adult criteria may be useful for older children and should be used when applicable.
E. Rheumatic fever, as used in this section assumes diagnosis made according to the revised Jones Criteria.

104.01 Category of Impairments, Cardiovascular
104.02 Chronic congestive failure. With two or more of the following signs:
A. Tachycardia (see Table I).
B. Tachypnea (see Table II).
C. Cardiomegaly on chest roentgenogram (see 104.00C).
D. Hepatomegaly (more than 2 cm. below the right costal margin in the right midclavicular line).
E. Evidence of pulmonary edema, such as rales or orthopnea.
F. Dependent edema.
G. Exercise intolerance manifested as laborored respiration on mild exertion (e.g., in an infant, feeding).

<table>
<thead>
<tr>
<th>TABLE I—TACHYCARDIA AT REST</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Age</strong></td>
</tr>
<tr>
<td>Under 1 yr</td>
</tr>
<tr>
<td>1 through 3 yrs</td>
</tr>
<tr>
<td>4 through 9 yrs</td>
</tr>
<tr>
<td>10 through 15 yrs</td>
</tr>
<tr>
<td>Over 15 yr</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>TABLE II—TACHYPIEKA AT REST</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Age</strong></td>
</tr>
<tr>
<td>Under 1 yr</td>
</tr>
<tr>
<td>1 through 5 yrs</td>
</tr>
<tr>
<td>6 through 9 yrs</td>
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<tr>
<td>Over 9 yrs</td>
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</tbody>
</table>

104.03 Hypertensive cardiovascular disease. With persistently elevated blood pressure for age (see Table III) and one of the following:
A. Impaired renal function as described under the criteria in 106.02; or
B. Cerebrovascular damage as described under the criteria in 111.06; or
C. Congestive heart failure as described under the criteria in 104.02.

**TABLE III—ELEVATED BLOOD PRESSURE**

<table>
<thead>
<tr>
<th>Age</th>
<th>S (over) mm.</th>
<th>Diastolic (over) in mm.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Under 6 mo</td>
<td>95</td>
<td>60</td>
</tr>
<tr>
<td>6 mo to 1 yr</td>
<td>110</td>
<td>70</td>
</tr>
<tr>
<td>1 through 8 yrs</td>
<td>115</td>
<td>80</td>
</tr>
<tr>
<td>9 through 11 yrs</td>
<td>120</td>
<td>80</td>
</tr>
<tr>
<td>12 through 15 yrs</td>
<td>130</td>
<td>80</td>
</tr>
<tr>
<td>Over 15 yrs</td>
<td>140</td>
<td>80</td>
</tr>
</tbody>
</table>

104.04 Cyanotic congenital heart disease. With one of the following:
A. Surgery is limited to palliative measures; or
B. Characteristic squatting, hemoptysis, syncope, or hypercyanotic spells; or
C. Chronic hematocrit of 55 percent or greater or arterial O₂ saturation of less than 95 percent at rest, or arterial oxygen tension of less than 60 Torr at rest.

104.05 Cardiac arrhythmia, such as persistent or recurrent heart block or A-V dissociation (with or without therapy). And one of the following:
A. Cardiac syncope; or
B. Congestive heart failure as described under the criteria in 104.02; or
C. Exercise intolerance with labored respirations on mild exertion (e.g., in infants, feeding).

104.07 Cardiac syncope with at least one documented syncopal episode characteristic of specific cardiac disease (e.g., aortic stenosis).

104.08 Recurrent hemoptysis. Associated with either pulmonary hypertension or extensive bronchial collaterals due to documented chronic cardiovascular disease.

104.09 Chronic rheumatic fever or rheumatic heart disease. With:
A. Persistence of rheumatic fever activity for 6 months or more, with significant murmur(s), cardiomegaly (see 104.00C), and other abnormal laboratory findings (such as elevated sedimentation rate or electrocardiographic findings); or
B. Congestive heart failure as described under the criteria in 104.02.

105.00 DIGESTIVE SYSTEM

A. Disorders of the digestive system which result in disability usually do so because of interference with nutrition and growth, multiple recurrent inflammatory lesions, or other complications of the disease. Such lesions or complications usually respond to treatment. To constitute a listed impairment, these must be shown to have persisted or be expected to persist despite prescribed therapy for a continuous period of at least 12 months.

B. Documentation of gastrointestinal impairments should include pertinent operative findings, radiographic studies, endoscopy, and biopsy reports. Where a liver biopsy has been performed in chronic liver disease, documentation should include the report of the biopsy.

C. Growth retardation and malnutrition. When the primary disorder of the digestive tract has been documented, evaluate resultant malnutrition under the criteria described in 105.08. Evaluate resultant growth impairment under the criteria described in 100.03. Intestinal disorders, including surgical diversions and potentially correctable congenital lesions, do not represent a severe impairment if the individual is able to maintain adequate nutrition growth and development.

D. Multiple congenital anomalies. See related criteria, and consider as a combination of impairments.

105.01 Category of Impairments, Digestive
105.03 Esophageal obstruction, caused by atresia, stricture, or stenosis with malnutrition as described under the criteria in 105.08.
105.05 Chronic liver disease. With one of the following:
A. Inoperable biliary atresia demonstrated by X-ray or surgery; or
B. Intractable ascites not attributable to other causes, with serum albumin of 3.0 gm./100 ml. or less; or
C. Esophageal varices (demonstrated by angiography, barium swallow, or endoscopy or by prior performance of a specific shunt or pllication procedure); or
D. Hepatic coma, documented by findings from hospital records; or
E. Hepatic encephalopathy. Evaluate under the criteria in 112.02; or
F. Chronic active inflammation or necrosis documented by SGOT persistently more than 100 units or serum bilirubin of 2.5 mg. percent or greater.

105.07 Chronic inflammatory bowel disease (such as ulcerative colitis, regional enteritis), as documented in 105.00. With one of the following:
A. Intestinal manifestations or complications, such as obstruction, abscess, or fistula formation which has lasted or is expected to last 12 months; or
B. Malnutrition as described under the criteria in 105.08; or
C. Growth impairment as described under the criteria in 100.02; or
D. Malnutrition, due to demonstrable gastrointestinal disease causing either a fall of 15 percentiles of weight which persists or the persistence of weight which is less than the third percentile (on standard growth charts). And one of the following:
A. Stool fat excretion per 24 hours:
   1. More than 15 percent in infants less than 6 months.
2. More than 10 percent in infants 6-18 months.
3. More than 6 percent in children more than 18 months; or
4. Persistent hematocrit of 30 percent or less despite prescribed therapy; or
5. Serum carotene of 40 mcg./100 ml. or less; or
6. Serum albumin of 3.0 gm./100 ml. or less.

106.00 GENITO-URINARY SYSTEM
A. Determination of the presence of chronic renal disease will be based upon the following factors:
   1. History, physical examination, and laboratory evidence of renal disease.

B. Renal transplant.

   The amount of function restored and the time required to affect improvement depend upon various factors including adequacy of post transplant renal function, incidence of renal infection, occurrence of rejection crisis, presence of systemic complications (anemia, neuropathy, etc.) and side effects of corticosteroid or immuno-suppressive agents. A period of at least 12 months is required for the individual to reach a point of stable medical improvement.

C. Evaluate associated disorders and complications according to the appropriate body system listing.

106.01 Category of Impairments, Genito-Urinary
106.02 Chronic renal disease. With:
   A. Persistent elevation of serum creatinine to 3 mg. per deciliter (100 ml.) or greater over at least 3 months; or
   B. Reduction of creatinine clearance to 30 ml. per minute (43 liters/24 hours) per 1.73 m² of body surface area over at least 3 months; or
   C. Chronic renal dialysis program for irreversible renal failure; or
   D. Renal transplant. Consider under a disability for 12 months following surgery; thereafter, evaluate the residual impairment (see 106.008).

106.04 Nephrotic syndrome, with edema not controlled by prescribed therapy. And:
   A. Serum albumin less than 2 gm./100 ml.; or
   B. Proteinuria more than 2.5 gm./1.73m²/day.

107.00 HEMIC AND LYMPHATIC SYSTEM
A. Sickle cell disease refers to a chronic hemolytic anemia associated with sickle cell hemoglobin, either homozygous or in combination with thalassemia or with another abnormal hemoglobin (such as C or F).

   Appropriate hematologic evidence for sickle cell disease, such as hemoglobin electrophoresis must be included. Vaso-occlusive, hemolytic, or aplastic episodes should be documented by description of severity, frequency, and duration.

   Disability due to sickle cell disease may be solely the result of a severe, persistent anemia or may be due to the combination of chronic progressive or episodic manifestations in the presence of a less severe anemia.

   Major visceral episodes causing disability include meningitis, osteomyelitis, pulmonary infections or infarctions, cerebrovascular accidents, congestive heart failure, genitourinary involvement, etc.

B. Coagulation defects. Chronic inherited coagulation disorders must be documented by appropriate laboratory evidence such as abnormal thromboplastin generation, coagulation time, or factor assay.

C. Acute leukemia. Initial diagnosis of acute leukemia must be based upon definitive bone marrow pathologic evidence. Recurrent disease may be documented by peripheral blood, bone marrow, or cerebrospinal fluid examination. The pathology report must be included.

   The designated duration of disability implicit in the finding of a listed impairment is contained in 107.11. Following the designated time period, a documented diagnosis itself is no longer sufficient to establish a severe impairment. The severity of any remaining impairment must be evaluated on the basis of the medical evidence.

107.01 Category of Impairments, Hemic and Lymphatic
107.02 Hemolytic anemia (due to any cause). Manifested by persistence of hematocrit of 26 percent or less despite prescribed therapy, and reticulocyte count of 4 percent or greater.

107.05 Sickle cell disease. With:
   A. Recent, recurrent, severe vaso-occlusive crises (musculoskeletal, vertebral, abdominal); or
   B. A major visceral complication in the 12 months prior to application; or
   C. A hyperhemolytic or aplastic crisis within 12 months prior to application; or
   D. Chronic, severe anemia with persistence of hematocrit of 26 percent or less; or
   E. Congestive heart failure, cerebrovascular damage, or emotional disorder as described under the criteria in 104.02, 111.00ff., or 152.00ff.

107.06 Chronic idiopathic thrombocytopenic purpura of childhood with purpura and thrombocytopenia of 40,000 platelets/cu. mm. or less despite prescribed therapy or recurrent upon withdrawal of treatment.

107.08 Inherited coagulation disorder. With:
   A. Repeated spontaneous or inappropriate bleeding; or
   B. Hemarthrosis with joint deformity.

107.11 Acute leukemia. Consider under a disability:
   A. For 2½ years from the time of initial diagnosis; or
B. For 2½ years from the time of recurrence of active disease.

109.00 [RESERVED]

109.00 ENDOCRINE SYSTEM

A. Cause of disability. Disability is caused by a disturbance in the regulation of the secretion or metabolism of one or more hormones which are not adequately controlled by therapy. Such disturbances or abnormalities usually respond to treatment. To constitute a listed impairment these must be shown to have persisted or be expected to persist despite prescribed therapy for a continuous period of at least 12 months.

B. Growth. Normal growth is usually a sensitive indicator of health as well as of adequate therapy in children. Impairment of growth may be disabling in itself or may be an indicator of a severe disorder involving the endocrine system or other body systems. Where involvement of other organ systems has occurred as a result of a primary endocrine disorder, these impairments should be evaluated according to the criteria under the appropriate sections.

C. Documentation. Description of characteristic history, physical findings, and diagnostic laboratory data must be included. Results of laboratory tests will be considered abnormal if outside the normal range or greater than two standard deviations from the mean of the testing laboratory. Reports in the file should contain the information provided by the testing laboratory as to their normal values for that test.

D. Hyperfunction of the adrenal cortex. Evidence of growth retardation must be documented as described in 100.00. Elevated blood or urinary free cortisol levels are not acceptable in lieu of urinary 17-hydroxycorticosteroid excretion for the diagnosis of adrenal cortical hyperfunction.

E. Adrenal cortical insufficiency. Documentation must include persistent low plasma cortisol or low urinary 17-hydroxycorticosteroids or 17-ketogenic steroids and evidence of unresponsiveness to ACTH stimulation.

109.01 Category of Impairments, Endocrine

109.02 Thyroid Disorders.

A. Hypothyroidism (as documented in 109.00C). With clinical manifestations despite prescribed therapy, and one of the following:

1. Elevated serum thyroxine (T4) and either elevated free T4 or resin T4 uptake; or
2. Elevated thyroid uptake of radiiodine; or
3. Elevated serum triiodothyronine (T3).

B. Hyperthyroidism. With one of the following, despite prescribed therapy:

1. IQ of 69 or less; or
2. Growth impairment as described under the criteria in 100.02A and B; or
3. Precocious puberty.

109.03 Hyperparathyroidism (as documented in 109.00C). With:

A. Repeated elevated total or ionized serum calcium; or

B. Elevated serum parathyroid hormone.

109.04 Hypoparathyroidism or Pseudohypoparathyroidism. With:

A. Severe recurrent tetany or convulsions which are unresponsive to prescribed therapy; or

B. Growth retardation as described under criteria in 100.02A and B.

109.05 Diabetes insipidus, documented by pathologic hypertonic saline or water deprivation test. And one of the following:

A. Intracranial space-occupying lesion, before or after surgery; or

B. Unresponsiveness to Pitressin; or

C. Growth retardation as described under the criteria in 100.02A and B; or

D. Unresponsive hypothalamic thirst center, with chronic or recurrent hypernatremia; or

E. Decreased visual fields attributable to a pituitary lesion.

109.06 Hyperfunction of the adrenal cortex (Primary or secondary). With:

A. Elevated urinary 17-hydroxycorticosteroids (or 17-ketogenic steroids) as documented in 109.00 C and D; and

B. Unresponsiveness to low-dose dexamethasone suppression.

109.07 Adrenal cortical insufficiency (as documented in 109.00 C and E) with recent, recurrent episodes of circulatory collapse.

109.08 Juvenile diabetes mellitus (as documented in 109.00C) requiring parenteral insulin. And one of the following, despite prescribed therapy:

A. Recent, recurrent hospitalizations with acidosis; or

B. Recent, recurrent episodes of hypoglycemia; or

C. Growth retardation as described under the criteria in 100.02A or B; or

D. Impaired renal function as described under the criteria in 106.00f.

109.09 Iatrogenic hypercorticoid state. With chronic glucocorticoid therapy resulting in one of the following:

A. Osteoporosis; or

B. Growth retardation as described under the criteria in 100.02A or B; or

C. Diabetes mellitus as described under the criteria in 100.08; or

D. Myopathy as described under the criteria in 111.06; or

E. Emotional disorder as described under the criteria in 112.00f.

109.10 Pituitary dwarfism (with documented growth hormone deficiency). And growth impairment as described under the criteria in 100.02B.

109.11 Adrenogenital syndrome. With:

A. Recent, recurrent self-losing episodes despite prescribed therapy; or
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B. Inadequate replacement therapy manifested by accelerated bone age and virilization, or
C. Growth impairment as described under the criteria in 110.02 A or B.

109.12 Hypoglycemia (as documented in 109.00 C). With recent, recurrent hypoglycemic episodes producing convulsion or coma.

109.13 Gonadal Dysgenesis (Turner’s Syndrome), chromosomally proven. Evaluate the resulting impairment under the criteria for the appropriate body system.

110.00 MULTIPLE BODY SYSTEMS

A. Catastrophic congenital abnormalities or disease. This section refers only to very serious congenital disorders, diagnosed in the newborn or infant child.

B. Immune deficiency diseases. Documentation of immune deficiency disease must be submitted, and may include quantitative immunoglobulins, skin tests for delayed hypersensitivity, lymphocyte stimulative tests, and measurements of cellular immunity mediators.

110.01 Category of Impairments, Multiple Body Systems

110.08 Catastrophic congenital abnormalities or disease. With:
A. A positive diagnosis (such as anencephaly, trisomy D or E, cyclopia, etc.), generally regarded as being incompatible with extrauterine life; or
B. A positive diagnosis (such as cri du chat, Tay-Sachs Disease) wherein attainment of the growth and development level of 2 years is not expected to occur.

110.09 Immune deficiency disease.
A. Hypogammaglobulinemia or dysgammaglobulinemia. With:
1. Recent, recurrent severe infections; or
2. A complication such as growth retardation, chronic lung disease, collagen disorder, or tumors.

E. Thymic dysplastic syndromes (such as Swiss, diGeorge).

111.00 NEUROLOGICAL

A. Seizure disorder must be substantiated by at least one detailed description of a typical seizure. Report of recent documentation should include an electroencephalogram and neurological examination. Sleep EEG is preferable, especially with temporal lobe seizures. Frequency of attacks and any associated phenomena should also be substantiated.

Young children may have convulsions in association with febrile illnesses. Proper use of 111.02 and 111.03 requires that a seizure disorder be established. Although this does not exclude consideration of seizures occurring during febrile illnesses, it does require documentation of seizures during nonfebrile periods.

There is an expected delay in control of seizures when treatment is started, particularly when changes in the treatment regimen are necessary. Therefore, a seizure disorder should not be considered to meet the requirements of 111.02 or 111.03 unless it is shown that seizures have persisted more than three months after prescribed therapy began.

B. Minor motor seizures. Classical petit mal seizures must be documented by characteristic EEG pattern, plus information as to age at onset and frequency of clinical seizures. Myoclonic seizures, whether of the typical infantile or Lennox-gastaut variety after infancy, must also be documented by the characteristic EEG pattern plus information as to age at onset and frequency of seizures.

C. Motor dysfunction. As described in 110.06, motor dysfunction may be due to any neurological disorder. It may be due to static or progressive conditions involving any area of the nervous system and producing any type of neurological impairment. This may include weakness, spasticity, lack of coordination, ataxia, tremor, athetosis, or sensory loss. Documentation of motor dysfunction must include neuologic findings and description of type of neurologic abnormality (e.g., spasticity, weakness), as well as a description of the child’s functional impairment (i.e., what the child is unable to do because of the abnormality). Where a diagnosis has been made, evidence should be included for substantiation of the diagnosis (e.g., blood chemistries and muscle biopsy reports), wherever applicable.

D. Impairment of communication. The documentation should include a description of a recent comprehensive evaluation, including all areas of affective and effective communication, performed by a qualified professional.

111.01 Category of Impairment, Neurological

111.02 Major motor seizure disorder.

A. Major motor seizures. In a child with an established seizure disorder, the occurrence of more than one major motor seizure per month despite at least three months of prescribed treatment. With:
1. Daytime episodes (loss of consciousness and convulsive seizures); or
2. Nocturnal episodes manifesting residuals which interfere with activity during the day.

B. Major motor seizures. In a child with an established seizure disorder, the occurrence of a least one major motor seizure in the year prior to application despite at least three months of prescribed treatment. And one of the following:
1. IQ of 69 or less; or
2. Significant interference with communication due to speech, hearing, or visual defect; or
3. Significant emotional disorder; or
4. Where significant adverse effects of medication interfere with major daily activities.

111.03 Minor motor seizure disorder. In a child with an established seizure disorder, the occurrence of more than one minor motor seizure per week, with alteration of awareness or loss of consciousness, despite at least three months of prescribed treatment.

111.05 Brain tumors. A. Malignant gliomas (astrocytoma—Grades III and IV, glioblastoma multiforme), medulloblastoma, ependymoblastoma, primary sarcoma or brain stem gliomas; or
B. Evaluate other brain tumors under the criteria for the resulting neurological impairment.

111.06 Motor dysfunction (due to any neurological disorder). Persistent disorganization or deficit of motor function for age involving two extremities, which (despite prescribed therapy) interferes with age-appropriate major daily activities and results in disruption of:
A. Fine and gross movements; or
B. Gait and station.

111.07 Cerebral palsy. With:
A. Motor dysfunction meeting the requirements of 111.06 or 101.03 or
B. Less severe motor dysfunction (but more than slight) and one of the following:
   1. IQ of 69 or less; or
   2. Seizure disorder, with at least one major motor seizure in the year prior to application; or
   3. Significant interference with communication due to speech, hearing or visual defect; or
   4. Significant emotional disorder.

111.08 Menigomyelocele (and related disorders). With one of the following despite prescribed treatment:
A. Motor dysfunction meeting the requirements of §101.03 or §111.06 or
B. Less severe motor dysfunction (but more than slight), and:
   1. Urinary or fecal incontinence when inappropriate for age; or
   2. IQ of 69 or less; or
   3. Four extremity involvement; or
   D. Noncompensated hydrocephalus producing interference with mental or motor developmental progression.

111.09 Communication impairment, associated with documented neurological disorder. And one of the following:
   A. Documented speech deficit which significantly affects the clarity and content of the speech; or
   B. Documented comprehension deficit resulting in ineffective verbal communication for age; or
   C. Impairment of hearing as described under the criteria in 102.08.

112.00 Mental and emotional disorders
A. Introduction. This section is intended primarily to describe mental and emotional disorders of young children. The criteria describing medically determinable impairments in adults should be used where they clearly appear to be more appropriate.
B. Mental retardation. General. As with any other impairment, the necessary evidence consists of symptoms, signs, and laboratory findings which provide medically demonstrable evidence of impairment severity. Standardized intelligence test results are essential to the adjudication of all cases of mental retardation that are not clearly covered under the provisions of 112.05A. Developmental milestone criteria may be the sole basis for adjudication only in cases where the child’s young age and/or condition preclude formal standardized testing by a psychologist or psychiatrist experienced in testing children.

Measures of intellectual functioning. Standardized intelligence tests, such as the Wechsler Preschool and Primary Scale of Intelligence (WPPSI), the Wechsler Intelligence Scale for Children—Revised (WISC-R), the Revised Stanford-Binet Scale, and the McCarthey Scales of Children’s Abilities, should be used wherever possible. Key data such as subtest scores should also be included in the report. Tests should be administered by a qualified and experienced psychologist or psychiatrist, and any discrepancies between formal tests results and the child’s customary behavior and daily activities should be duly noted and resolved.

Developmental milestone criteria. In the event that a child’s young age and/or condition preclude formal testing by a psychologist or psychiatrist experienced in testing children, a comprehensive evaluation covering the full range of developmental activities should be performed. This should consist of a detailed account of the child’s daily activities together with direct observations by a professional person; the latter should include indices or manifestations of social, intellectual, adaptive, verbal, motor (posture, locomotion, manipulation), language, emotional, and self-care development for age. The above should then be related by the evaluating or treating physician to established developmental norms of the kind found in any widely used standard pediatrics text.
C. Profound combined mental-neurological-musculoskeletal impairments. There are children with profound and irreversible brain damage resulting in total incapacitation. Such children may meet criteria in either neurological, musculoskeletal, and/or mental sections; they should be adjudicated under the criteria most completely substantiated by the medical evidence submitted. Frequently, the most appropriate criteria will
be found under the mental impairment section.

112.01 Category of Impairments, Mental and Emotional

112.02 Chronic brain syndrome. With arrest of developmental progression for at least six months or loss of previously acquired abilities; or

112.03 Psychosis of infancy and childhood. Documented by psychiatric evaluation and supported, if necessary, by the results of appropriate standardized psychological tests and manifested by marked restriction in the performance of daily age-appropriate activities; constriction of age-appropriate interests; deficiency of age-appropriate self-care skills; and impaired ability to relate to others; together with persistence of one (or more) of the following:

A. Significant withdrawal or detachment; or
B. Impaired sense of reality; or
C. Bizarre behavior patterns; or
D. Strong need for maintenance of sameness, with intense anxiety, fear, or anger when change is introduced; or
E. Panic at threat of separation from parent.

112.04 Functional nonpsychotic disorders. Documented by psychiatric evaluation and supported, if necessary, by the results of appropriate standardized psychological tests and manifested by marked restriction in the performance of daily age-appropriate activities; constriction of age-appropriate interests; deficiency of age-appropriate self-care skills; and impaired ability to relate to others; together with persistence of one (or more) of the following:

A. Psychophysiological disorder (e.g., diarrhea, asthma); or
B. Anxiety; or
C. Depression; or
D. Phobic, obsessive, or compulsive behavior; or
E. Hypochondriasis; or
F. Hysteria; or
G. Asocial or antisocial behavior.

112.05 Mental retardation. Achievement of only those developmental milestones generally acquired by children no more than one-half the child’s chronological age; or

B. IQ of 59 or less; or
C. IQ of 60-69, inclusive, and a physical or other mental impairment imposing additional and significant restriction of function or developmental progress.

113.00 NEOPLASTIC DISEASES, MALIGNANT

A. Introduction. Determination of disability in the growing and developing child with a malignant neoplastic disease is based upon the combined effects of:

1. The pathophysiology, histology, and natural history of the tumor; and
2. The effects of the currently employed aggressive multimodal therapeutic regimens.

Combinations of surgery, radiation, and chemotherapy or prolonged therapeutic schedules impart significant additional morbidity to the child during the period of greatest risk from the tumor itself. This period of highest risk and greatest therapeutically-induced morbidity defines the limits of disability for most of childhood neoplastic disease.

B. Documentation. The diagnosis of neoplasm should be established on the basis of symptoms, signs, and laboratory findings. The site of the primary, recurrent, and metastatic lesion must be specified in all cases of malignant neoplastic diseases. If an operative procedure has been performed, the evidence should include a copy of the operative note and the report of the gross and microscopic examination of the surgical specimen, along with all pertinent laboratory and X-ray reports. The evidence should also include a recent report directed especially at describing whether there is evidence of local or regional recurrence, soft part or skeletal metastases, and significant post therapeutic residuals.

C. Malignant solid tumors, as listed under 113.03, include the histiocytosis syndromes except for solitary eosinophilic granuloma. Thus, 113.03 should not be used for evaluating brain tumors (see 111.05) or thyroid tumors, which must be evaluated on the basis of whether they are controlled by prescribed therapy.

D. Duration of disability from malignant neoplastic tumors is included in 113.02 and 113.03. Following the time periods designated in these sections, a documented diagnosis itself is no longer sufficient to establish a severe impairment. The severity of a remaining impairment must be evaluated on the basis of the medical evidence.

113.01 Category of Impairments, Neoplastic Diseases—Malignant

113.02 Lymphoreticular malignant neoplasms. A. Hodgkin’s disease with progressive disease not controlled by prescribed therapy; or
B. Non-Hodgkin’s lymphoma. Consider under a disability:

1. For 2½ years from time of initial diagnosis; or
2. For 2½ years from time of recurrence of active disease.

113.03 Malignant solid tumors. Consider under a disability:

A. For 2 years from the time of initial diagnosis; or
B. For 2 years from the time of recurrence of active disease.

113.04 Neuroblastoma. With one of the following:

A. Extension across the midline; or
B. Distant metastases; or
C. Recurrence; or
D. Onset at age 1 year or older.
APPENDIX 2 TO PART 220—MEDICAL-VOCATIONAL GUIDELINES

Sec.
200.00 Introduction.
201.00 Maximum sustained work capability limited to sedentary work as a result of severe medically determinable impairment(s).
202.00 Maximum sustained work capability limited to light work as a result of severe medically determinable impairment(s).
203.00 Maximum sustained work capability limited to medium work as a result of severe medically determinable impairment(s).
204.00 Maximum sustained work capability limited to heavy work (or very heavy work) as a result of severe medically determinable impairment(s).
200.00 Introduction. (a) The following rules reflect the major functional and vocational patterns which are encountered in cases which cannot be evaluated on medical considerations alone, where an individual with a severe medically determinable physical or mental impairment(s) is not engaging in substantial gainful activity and the individual’s impairment(s) prevents the performance of his or her vocationally relevant past work. They also reflect the analysis of the various vocational factors (i.e., age, education, and work experience) in combination with the individual’s residual functional capacity (used to determine his or her maximum sustained work capability for sedentary, light, medium, heavy, or very heavy work) in evaluating the individual’s ability to engage in substantial gainful activity in other than his or her vocationally relevant past work. Where the findings of fact made with respect to a particular individual’s vocational factors and residual functional capacity coincide with all of the criteria of a particular rule, the rule directs a conclusion as to whether the individual is or is not disabled. However, each of these findings of fact is subject to rebuttal and the individual may present evidence to refute such findings. Where any one of the findings of fact does not coincide with the corresponding criterion of a rule, the rule does not apply in that particular case and, accordingly, does not direct a conclusion of disabled or not disabled. In any instance where a rule does not apply, full consideration must be given to all of the relevant facts of the case in accordance with the definitions and discussions of each factor in the appropriate sections of the regulations.
(b) The existence of jobs in the national economy is reflected in the “Decisions” shown in the rules; i.e., in promulgating the rules, administrative notice has been taken of the numbers of unskilled jobs that exist throughout the national economy at the various functional levels (sedentary, light, medium, heavy, and very heavy) as supported by the “Dictionary of Occupational Titles” and the “Occupational Outlook Handbook,” published by the Department of Labor; the “County Business Patterns” and “Census Surveys” published by the Bureau of the Census; and occupational surveys of light and sedentary jobs prepared for the Social Security Administration by various State employment agencies. Thus, when all factors coincide with the criteria of a rule, the existence of such jobs is established. However, the existence of such jobs for individuals whose remaining functional capacity or other factors do not coincide with the criteria of a rule must be further considered in terms of what kinds of jobs or types of work may be either additionally indicated or precluded.
(c) In the application of the rules, the individual’s residual functional capacity (i.e., the maximum degree to which the individual retains the capacity for sustained performance of the physical-mental requirements of jobs), age, education, and work experience must first be determined.
(d) The correct disability decision (i.e., on the issue of ability to engage in substantial gainful activity) is found by then locating the individual’s specific vocational profile. If an individual’s specific profile is not listed within this appendix 2, a conclusion of disabled or not disabled is not directed. Thus, for example, an individual’s ability to engage in substantial gainful work where his or her residual functional capacity falls between the ranges of work indicated in the rules (e.g., the individual who can perform more than light but less than medium work), is decided on the basis of the principles and definitions in the regulations, giving consideration to the rules for specific case situations in this appendix 2. These rules represent various combinations of exertional capabilities, age, education and work experience and also provide an overall structure for evaluation of those cases in which the judgments as to each factor do not coincide with those of any specific rule. Thus, when the necessary judgments have been made as to each factor and it is found that no specific rule applies, the rules still provide guidance for decisionmaking, such as in cases involving combinations of impairments. For example, if strength limitations resulting from an individual’s impairment(s) considered with the judgments made as to the individual’s

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113.05 Retinoblastoma. With one of the following:
A. Bilateral involvement; or
B. Metastases; or
C. Extension beyond the orbit; or
D. Recurrence.