

To minimize disruption to the MDM coding portion of the encounter, the American Medical Association (AMA) has revised and updated the office or other outpatient E/M services MDM terminology and their definitions. Understanding the meaning of these terms and their definitions will assist the dermatologist or non-physician clinician (NPC) in consistently and accurately applying the MDM concepts when selecting the level of service for dermatology encounters.

The examples provided do not constitute an exhaustive list. **Conditions listed can qualify under multiple categories, depending on disease severity and patient presenting circumstances at the time of the encounter.**

Terminology	Definition
Problem	<p>A problem is a disease, condition, illness, injury, symptom, sign, finding, complaint, or other matter addressed at the encounter, with or without a diagnosis being established at the time of the encounter.</p> <p><i>Examples include any problem presented and addressed during an encounter including but not limited to: Acne, Actinic Keratosis, Allergic contact dermatitis, Alopecia, Atopic Dermatitis, Basal Cell Carcinoma (BCC), Irritant Dermatitis, Seborrheic Keratosis, Squamous Cell Carcinoma (SCC), Neoplasm Uncertain Behavior (NUB), Pigmented lesion, Pruritus, Psoriasis, Rash, etc.</i></p>
Problem addressed	<p>A problem is addressed or managed when it is evaluated or treated at the encounter by the dermatologist or NPC reporting the service.</p> <p><i>This includes consideration or lack thereof of further testing or treatment that may not be elected by virtue of risk/benefit analysis or the patient/parent/guardian/surrogate's choice.</i></p> <p><i>Notation in the patient's medical record that another professional is managing the problem without additional assessment or care coordination documented does not qualify as being 'addressed' or managed by the dermatologist or NPC reporting the service.</i></p> <p><i>Referral without evaluation (by history, exam, or diagnostic study[ies]) or consideration of treatment does not qualify as being addressed or managed.</i></p>
Minimal problem	<p>A problem that may not require the presence of the dermatologist or NPC, but the service is provided under the supervision of a dermatologist or NPC.</p> <p><i>Examples include Suture removal during a nurse encounter (see 99211).</i></p>

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<p>Self-limited or minor problem</p>	<p>A problem that runs a definite and prescribed course, is transient in nature, and is not likely to permanently alter health status.</p> <p><i>Examples may include:</i> <i>Acrochordon, Angiofibroma, Bug bite, Dry skin or Xerosis, Epidermal inclusion cyst, Lentiginos, Lipoma, Melanocytic nevi, Post Inflammatory Hyperpigmentation, Rhytids, Scar, Seborrhic Keratosis, Telangiectasi, etc.</i></p>
<p>Stable, chronic illness</p>	<p>A problem with an expected duration of at least a year or until the death of the patient. For the purpose of defining chronicity, conditions are treated as chronic whether or not stage or severity changes.</p> <p>'Stable' for the purposes of categorizing medical decision making is defined by the specific treatment goals for an individual patient. A patient that is not at their treatment goal is not stable, even if the condition has not changed and there is no short-term threat to life or function.</p> <p><i>Examples, when stable, may include:</i> <i>Acne, Actinic Keratoses, Actinic Cheilitis, Alopecia Areata, Atopic Dermatitis, Atypical Nevi, Bullous Pemphigoid, Chronic Urticaria, Dermatomyositis, Diffuse Actinic Damage, Disseminated superficial actinic porokeratosis (DSAP), Granuloma Annulare, History of skin cancer, Intertrigo, Keloid, Melasma, Notalgia Paresthetica, Onychomycosis, uncomplicated, Pemphigus Vulgaris, Psoriasis Vulgaris, Rosacea, Scleroderma, Seborrhic Dermatitis, Sjogren Syndrome, Stasis ulcer, etc.</i></p>
<p>Acute, uncomplicated illness or injury</p>	<p>A recent or new short-term problem with low risk of morbidity for which treatment is considered. There is little to no risk of mortality with treatment, and full recovery without functional impairment is expected.</p> <p>A problem that is normally self-limited or minor but is not resolving consistent with a definite and prescribed course is an acute uncomplicated illness.</p> <p><i>Examples may include:</i> <i>Abrasion, Acne, Actinic Keratosis, Acute Urticaria, Allergic Contact Dermatitis, Atypical Nevus, Cellulitis, Epidermal cyst -inflamed, Erythema nodosum -uncomplicated, Folliculitis, Impetigo, Inflamed or infected Epidermal Inclusion Cyst, Inflamed Seborrhic Keratosis, Irritant Dermatitis, Intertrigo, Melanoma in-situ, Melasma during pregnancy, NCCN Low risk NMSC, NCCN low risk BCC/SCC, Onychomycosis -uncomplicated, Tinea Corporis, Retinoid Dermatitis, Squamous Cell Carcinoma In-Situ (SCCIS), Wound healing by second intent, etc.</i></p>
<p>Chronic illness with exacerbation, progression, or side effects of treatment</p>	<p>A chronic illness that is acutely worsening, poorly controlled or progressing with an intent to control progression and requiring additional supportive care or requiring attention to treatment for side effects, but that does not require consideration of hospital level of care.</p> <p><i>Examples may include:</i> <i>Acne -flaring, Alopecia areata, Atopic Dermatitis -flaring, Discoid Lupus, Erythematosus (DLE) with new, active lesions, Drug Eruption, Immunocompromised/ Immunosuppressed patient with skin cancer, Onychomycosis -uncomplicated, Psoriasis that has spread to other anatomic location(s), Psoriasis vulgaris with new flare, poor progression or not at treatment goal, Subacute lupus -flaring, etc.</i></p>

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<p>Undiagnosed new problem with uncertain prognosis</p>	<p>A problem in the differential diagnosis that represents a condition likely to result in a high risk of morbidity without treatment.</p> <p><i>An example may include: Changing pigmented lesion, New bleeding red papule, Atypical Fibroxanthoma (AFX), NCCN Intermediate Risk SCC, NCCN High Risk BCC / SCC, Area H NMSC, Melanoma in situ, Atypical Nevus, Neoplasm of Uncertain Behavior (NUB), T1a Invasive Malignant Melanoma, Rash, Leg ulcer, etc.</i></p>
<p>Acute illness with systemic symptoms</p>	<p>An illness that causes systemic symptoms and has a high risk of morbidity without treatment.</p> <p>For systemic general symptoms such as fever, body aches, or fatigue in a minor illness that may be treated to alleviate symptoms, shorten the course of illness, or to prevent complications, see the definitions for 'self-limited or minor' or 'acute, uncomplicated.' Systemic symptoms may not be general but may be single system.</p> <p><i>Examples may include: New onset acute Systemic lupus erythematosus, Cellulitis with fever and chills, Drug-induced exfoliative erythroderma, Erythema multiforme, Leukocytoclastic vasculitis with hematuria, Pemphigus vulgaris -flaring, Psoriasis with psoriatic arthritis, Tick bite with myalgias, Viral exanthema with systemic symptoms, Varicella Zoster with neuralgia, Worsening pyoderma gangrenosum with abdominal symptoms</i></p>
<p>Acute, complicated injury</p>	<p>An injury which requires treatment that includes evaluation of body systems that are not directly part of the injured organ, the injury is extensive, or the treatment options are multiple and/or associated with risk of morbidity.</p> <p><i>Examples may include: Severe fall in the examination room with head trauma, Severe, extensive blistering (second degree) sunburn</i></p>
<p>Chronic illness with severe exacerbation, progression, or side effects of treatment</p>	<p>The severe exacerbation or progression of a chronic illness or severe side effects of treatment that have significant risk of morbidity and may require hospital level of care.</p> <p><i>Examples may include: Acne fulminans flare after initiation of isotretinoin therapy, Pemphigus vulgaris with severe cutaneous and oral mucosal/esophageal exacerbation, Dermatomyositis with worsening muscle weakness, Systemic lupus erythematosus with acute diffuse purpuric eruption, Erythrodermic psoriasis with systemic symptoms</i></p>
<p>Acute or chronic illness or injury that poses a threat to life or bodily function</p>	<p>An acute illness with systemic symptoms, or an acute complicated injury, or a chronic illness or injury with exacerbation and/or progression or side effects of treatment, that poses a threat to life or bodily function in the near term without treatment.</p> <p><i>Examples may include: Advanced regional Melanoma, AJCC8 T3/T4 SCC, BWH T2B/T3 SCC, Drug Reaction with Eosinophilia and Systemic Symptoms (DRESS), Erythroderma with hypotension, Sezary Syndrome, Merkel Cell Carcinoma, Metastatic SCC or Melanoma, New invasive T1b or higher T Melanoma, Paraneoplastic pemphigus, Toxic epidermal necrolysis</i></p>

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Test

Tests are imaging, laboratory, psychometric, or physiologic data. A clinical laboratory panel (e.g., basic metabolic panel [80047]) is a single test.

For the purposes of data reviewed and analyzed, pulse oximetry is not a test.

The determination between single or multiple unique tests is defined by the CPT code. Defined panels of tests, e.g., Comprehensive Metabolic Panel (CMP), Complete Blood Count (CBC) each count as one individual test.

Cholesterol (82465), triglycerides (84478) and quantitative human chorionic gonadotropin (hCG) (84702) would each be counted as individual tests when they are not performed as part of the lipid panel.

Examples may include review of:

Skin biopsy report not generated by the treating practitioner, CBC, differential, platelet, CMP, Chest X-ray, LDH level when ordered as a single test, CT scan, MRI, Lipid panel, Pregnancy test, Prothrombin Time Test/International, Normalized Ratio (PT/INR), Medication Blood Level

Ordering a test(s) includes both the order and the analysis of the test result. As such, the review of the ordered test result(s) is part of the encounter at which the test is ordered and is counted only once under data reviewed element.

Tests ordered are presumed to be analyzed when the results are reported, even if the analysis is performed post-encounter. Therefore, when they are ordered during an encounter, they are counted as part of that encounter.

Ordering a test may include those considered, but not selected after shared decision making, such as NOT ordering a chest X-ray for a melanoma diagnosis.

All considerations must be documented in the medical record including tests that may normally be performed, but due to the risk for a specific patient are not ordered.

Any service for which the professional component is separately reported by the physician or other QHP reporting the E/M services is not counted as a data element ordered, reviewed, analyzed, or independently interpreted for the purposes of determining the level of MDM.

An example may include when a dermatologist or QHP reports the pathology code 88305. The ordering and reviewing of the histopathology report cannot be counted toward the E/M data element.

Tests that are ordered outside of an encounter may be counted as part of the data element during the encounter in which they are analyzed.

This may include analyzing the results of the tests that are ordered outside of the face-to-face encounter.

When the ordering of the test does not occur during an E/M encounter, analyzing the test results can be counted as a single test during the encounter at which the result is reviewed.

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	<p>When multiple results of the same unique test (e.g., serial blood glucose values) are compared during an E/M service, count it as one unique test.</p> <p><i>For example, an encounter that includes an order for monthly prothrombin times would count for one prothrombin time ordered and reviewed.</i></p>
	<p>Tests that have overlapping elements are not unique, even if some of their individual components are identified with distinct CPT codes.</p> <p><i>For example, a CBC with differential would incorporate the set of hemoglobin, CBC without differential, and platelet count.</i></p>
Analyzed	<p>Is the process of using test data as part of the MDM.</p> <p><i>The data element itself may not be subject to analysis (e.g., glucose), but it is instead included in the thought processes for diagnosis, evaluation, or treatment</i></p>
Unique Source	<p>A unique source is defined as a physician or QHP in a distinct group or different specialty or subspecialty, or a unique entity.</p> <p>Please check directly with the private payer and seek clarification on how they will apply this policy.</p> <p><i>Review of all materials from any unique source counts as one element toward MDM.</i></p>
Combination of Data Elements	<p>A combination of different data elements does not require each item type or category to be represented.</p> <p>A unique test ordered, plus a note reviewed from an external source and an independent historian would be a combination of three elements.</p> <p><i>This may include a combination of notes reviewed, tests ordered, tests reviewed, or independent historian, which allows these elements to be summed: External records, Skin biopsy result, PT/INR, Caregiver/Power of Attorney/Historian/Parent/Guardian/Witness, Test result and/or Independent interpretation of test</i></p>
External	<p>External records, communications, and/or test results are from an external physicians, NPC, facility, or health care organization not affiliated with the practice.</p> <p><i>Data to be reviewed can also include information obtained from multiple sources or interprofessional communication that is not separately reported e.g., medical records, tests, and/or other information that must be obtained, ordered, reviewed, and analyzed for the encounter.</i></p>

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<p>External physician or other qualified health care professional</p>	<p>An external physician or other qualified health care professional is an individual who is not in the same group practice or is of a different specialty or subspecialty.</p> <p>It includes licensed professionals that are practicing independently. It may also be a facility or organizational provider such as in a hospital, nursing facility, or home health care agency.</p> <p>Though Medicare recognizes sub-specialty credentialing, private payers do not.</p> <p>Please check directly with the private payer and seek clarification on how they will apply this policy.</p> <p><i>This may include review of external records, communications, and/or test results from a physician, NPC, facility, or health care organization not affiliated with the dermatology practice or are from a different specialty or subspecialty.</i></p> <p><i>Discussion requires an interactive exchange. The exchange must be direct and not through intermediaries (e.g., clinical staff or trainees).</i></p> <p><i>Sending chart notes or written exchanges that are within progress notes does not qualify as an interactive exchange.</i></p> <p><i>The discussion does not need to be on the date of the encounter, but it is counted only once and only when it is used in the decision making of the encounter.</i></p> <p><i>It may be asynchronous (i.e., does not need to be in person), but it must be initiated and completed within a short time period (e.g., within a day or two).</i></p> <p><i>Examples may include:</i> <i>Referral or consult from any specialty other than dermatology within the same practice group, Referral from any outside group, including dermatology</i></p>
<p>Independent historian(s)</p>	<p>An individual, other than the patient or physician (e.g., parent, guardian, surrogate, spouse, witness) who provides a history in addition to a history provided by the patient who is unable to provide a complete or reliable history (e.g., due to developmental stage, dementia, or psychosis) or because the dermatologist or NPC determines that a confirmatory history is judged to be necessary.</p> <p>In the case where there may be conflict or poor communication between multiple historians and more than one historian(s) is needed, the independent historian(s) requirement is met.</p> <p>The independent history does not need to be obtained in person but does need to be obtained directly from the historian providing the independent information.</p> <p><i>Translators are not considered an independent historian, as they only translate the patient words and are not adding to the history being obtained.</i></p>

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Independent interpretation	<p>The interpretation of a test for which there is a CPT code, and an interpretation or report is customary. This does not apply when the dermatologist or NPC is reporting the service or has previously reported the service for the patient.</p> <p>Documentation of the interpretation test result must be documented in the patient medical record.</p> <p><i>This includes the interpretation and/or reporting of results of tests not ordered by the dermatologist or NPC, review of slides as part of a request for consultation by another physician.</i></p> <ul style="list-style-type: none">• <i>Independent assessment of pathology slides from an external referral with your own interpretation documented in the chart</i>• <i>Review of a CT scan or MRI (the images) prior to performing surgery with your own interpretation documented in the chart</i>
Appropriate source	<p>For the purpose of the Discussion of Management data element, an appropriate source includes professionals who are not health care professionals but may be involved in the management of the patient).</p> <p>It does not include discussion with family or informal caregivers.</p> <p><i>This includes lawyer, parole officer, case manager, teacher</i></p>
Risk	<p>One element used in selecting the level of service is the risk of complications and/or morbidity or mortality of patient management at an encounter. This is distinct from the risk of the condition itself.</p> <p>The probability and/or consequences of an event. The assessment of the level of risk is affected by the nature of the event under consideration. For example, a low probability of death may be high risk, whereas a high chance of a minor, self-limited adverse effect of treatment may be low risk.</p> <p>Definitions of risk are based upon the usual behavior and thought processes of a physician or other qualified health care professional in the same specialty. Trained clinicians apply common language usage meanings to terms such as 'high,' 'medium,' 'low,' or 'minimal' risk and do not require quantification for these definitions, (though quantification may be provided when evidence-based medicine has established probabilities).</p> <p><i>The risk of patient management criteria applies to the patient management decisions made by the reporting physician or other QHP as part of the reported encounter.</i></p> <p><i>This may include patient management decisions made during the visit, associated with the patient's problem(s), the diagnostic procedure(s), treatment(s).</i></p> <p><i>It also includes the possible management options selected and those considered but not selected, after sharing the MDM with the patient and/or family.</i></p>

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Morbidity	<p>A state of illness or functional impairment that is expected to be of substantial duration during which function is limited, quality of life is impaired, or there is organ damage that may not be transient despite treatment.</p> <p>For the purposes of medical decision making, level of risk is based upon consequences of the problem(s) addressed at the encounter when appropriately treated.</p> <p>Risk also includes medical decision making related to the need to initiate or forego further testing, treatment, and/or hospitalization.</p>
Social Determinants of Health (SDOH)	<p>Economic and social conditions that influence the health of people and communities.</p> <p><i>Examples may include:</i> <i>food or housing insecurity, lack of reliable transportation to medical appointments, homelessness, financial insecurity, etc.</i></p>
Drug therapy requiring intensive monitoring	<p>A drug that requires intensive monitoring is a therapeutic agent that has the potential to cause serious morbidity or death. The monitoring is performed for assessment of these adverse effects and not primarily for assessment of therapeutic efficacy.</p> <p>The monitoring should be that which is generally accepted practice for the agent but may be patient specific in some cases. Intensive monitoring may be long-term or short term.</p> <p>Long-term intensive monitoring is not less than quarterly. The monitoring may be by a lab test, a physiologic monitoring for test, or imaging. Monitoring by history or examination does not qualify. The monitoring affects the level of toxicity medical decision making in an encounter in which it is considered in the management of the patient.</p> <p><i>This may include:</i></p> <ul style="list-style-type: none">• <i>Psoriasis patient on methotrexate or cyclosporine with labs drawn and analyzed four times per year</i>• <i>Patient with elevated lipids or an oral retinoid requiring monitoring four times per year or more</i>• <i>Pemphigus patient treated with cyclophosphamide</i> <p><i>Quantitative human chorionic gonadotropin (hCG) levels during isotretinoin therapy does NOT qualify</i></p>

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**Surgery
(minor or major,
elective, emergency,
procedure,
or patient risk)**

Minor or Major:

The classification of surgery into minor or major is based on the common meaning of such terms when used by trained clinicians, similar to the use of the term “risk.” These terms are not defined by a surgical package classification.

Elective or Emergency:

Elective procedures and emergent or urgent procedures describe the timing of a procedure when the timing is related to the patient’s condition. An elective procedure is typically planned in advance (e.g., scheduled for weeks later), while an emergent procedure is typically performed immediately or with minimal delay to allow for patient stabilization. Both elective and emergent procedures may be minor or major procedures.

Risk factors are those that are relevant to the patient and procedure. Evidence-based risk calculators may be used, but are not required, in assessing patient and procedure risk.

Minor surgeries can include:

Skin biopsy, Excision, Destruction, Shave removal, Linear closure, Uncomplicated Mohs surgery

Major surgeries can include:

Complex or extensive excision, Complex Mohs surgery, Flap reconstruction, Graft reconstruction

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