STATE OF MICHIGAN STATE OFFICE OF ADMINISTRATIVE HEARINGS AND RULES FOR THE DEPARTMENT OF COMMUNITY HEALTH

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IN THE MAT	TER OF:	Docket No. 2011 Case No. 382	
Appell	ant.		
	DECISION A	ND ORDER	
	s before the undersigned Administration CFR 431.200 et seq., following		
After due not behalf. Her	ice, a hearing was held , appea	. The Appellant appeared and testified on her behalf	
	opeared on behalf of acted Medicaid provider. appeared on behalf	, a Department of of the plan.	for for
ISSUE			
	e Department properly deny the policy osteogenesis bone stimulate		request for
FINDINGS O	F FACT		
	trative Law Judge (ALJ), based of the whole record, finds as materi		d substantia
1.	The Appellant is a Medicaid be	eneficiary enrolled with	
2.	The Appellant has had a broke has not healed despite medical	•	. The toe
3.	The Appellant's physician so osteogenesis bone stimulator to	•	n ultrasonio
4.	equipment and denied the req	reviewed the reques uest on , o	ted medica citing interna

5. On Rules received a hearing request.

CONCLUSIONS OF LAW

The Medical Assistance Program is established pursuant to Title XIX of the Social Security Act and is implemented by Title 42 of the Code of Federal Regulations (CFR). It is administered in accordance with state statute, the Social Welfare Act, the Administrative Code, and the State Plan under Title XIX of the Social Security Act Medical Assistance Program.

On May 30, 1997, the Department received approval from the Health Care Financing Administration, U.S. Department of Health and Human Services, allowing Michigan to restrict Medicaid beneficiaries' choice to obtain medical services only from specified Medicaid Health Plans.

The Respondent is one of those Medicaid Health Plans.

The covered services that the Contractor has available for enrollees must include, at a minimum, the covered services listed below (List omitted by Administrative Law Judge). The Contractor may limit services to those which are medically necessary and appropriate, and which conform to professionally accepted standards of care. The Contractor must operate consistent with all applicable Medicaid provider manuals and publications for coverages and limitations. If new services are added to the Michigan Medicaid Program, or if services are expanded, eliminated, or otherwise changed, the Contractor must implement the changes consistent with State direction in accordance with the provisions of Contract Section 2.024.

Section 1.022(E)(1), Covered Services. MDCH contract (Contract) with the Medicaid Health Plans, October 1, 2009.

- (1) The major components of the Contractor's utilization management (UM) program must encompass, at a minimum, the following:
 - (a) Written policies with review decision criteria and procedures that conform to managed health care industry standards and processes.

- (b) A formal utilization review committee directed by the Contractor's medical director to oversee the utilization review process.
- (c) Sufficient resources to regularly review the effectiveness of the utilization review process and to make changes to the process as needed.
- (d) An annual review and reporting of utilization review activities and outcomes/interventions from the review.
- (e) The UM activities of the Contractor must be integrated with the Contractor's QAPI program.

(2) Prior Approval Policy and Procedure

The Contractor must establish and use a written prior approval policy and procedure for UM purposes. The Contractor may not use such policies and procedures to avoid providing medically necessary services within the coverages established under the Contract. The policy must ensure that the review criteria for authorization decisions are applied consistently and require that the reviewer consult with the requesting provider when appropriate. The policy must also require that UM decisions be made by a health care professional who has appropriate clinical expertise regarding the service under review.

Section 1.022(AA)(1) and (2), Utilization Management, Contract, October 1, 2009.

As it says in the above Department - MHP contract language, a MHP such as may limit services to those that are medically necessary and that are consistent with applicable Medicaid Provider Manuals. It may require prior authorization for certain procedures. The process must be consistent with the Medicaid Provider Manual. The pertinent section of the Medicaid Provider Manual criteria for Medical Necessity is below.

The Medicaid Provider Manual provides, in pertinent part, as follows:

1.5 MEDICAL NECESSITY

Medical devices are covered if they are the most costeffective treatment available and meet the Standards of Coverage stated in the Coverage Conditions and Requirements Section of this chapter. The medical record must contain sufficient documentation of the beneficiary's medical condition to substantiate the necessity for the type

and quantity of items ordered and for the frequency of use or replacement. The information should include beneficiary's diagnosis, medical condition, and other pertinent information including, but not limited to, duration of the condition, clinical course, prognosis, nature and extent of functional limitations, other therapeutic interventions and results, and past experience with related items. Neither a physician's order nor a certificate of medical necessity by itself provides sufficient documentation of medical necessity. even though it is signed by the treating physician. Information in the medical record must support the item's medical necessity and substantiate that the medical device needed is the most appropriate economic alternative that meets MDCH standards of coverage. Medical equipment may be determined to be medically necessary when all of the following apply:

- Within applicable federal and state laws, rules, regulations, and MDCH promulgated policies.
- Medically appropriate and necessary to treat a specific medical diagnosis or medical condition, or functional need, and is an integral part of the nursing facility daily plan of care or is required for the community residential setting.
- Within accepted medical standards; practice guidelines related to type, frequency, and duration of treatment; and within scope of current medical practice.
- Inappropriate to use a nonmedical item.
- The most cost effective treatment available.
- It is ordered by the treating physician, and clinical documentation from the medical record supports the medical necessity for the request (as described above) and substantiates the physician's order.
- It meets the standards of coverage published by MDCH.
- It meets the definition of Durable Medical Equipment (DME), as defined in the Program Overview section of this chapter.
- Its use meets FDA and manufacturer indications.

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1.10 NONCOVERED ITEMS

Items that are not covered by Medicaid include, but are not limited to:

- Adaptive equipment (e.g., rocker knife, swivel spoon, etc.)
- Air conditioner
- Air purifier
- Devices used for play, pre-mobility development, or exercise are not considered pediatric mobility devices for the purpose of reimbursement and are not covered (e.g., jet mobile, ready racer, creepster crawler)
- Enteral formula to accommodate psychological or behavioral conditions, food preferences, allergies, loss of appetite, or noncompliance with a specialized diet
- Environmental Control Units
- Equipment not used or not used properly by the beneficiary
- Equipment for social or recreational purposes
- Exam tables/massage tables
- Exercise equipment (e.g., tricycles, exercise bikes, weights, mat/mat tables, etc.)
- Generators
- Hand/body wash
- Heating pads
- Home modifications
- Hot tubs
- House/room humidifier
- Ice packs
- Items for a beneficiary who is non-compliant with a physician's plan of care (or) items ordered for the purpose of solving problems related to noncompliance (e.g., insulin pump)
- Items used solely for the purpose of restraining the beneficiary for behavioral or other reasons
- Lift chairs, reclining chairs, vibrating chairs
- More than one pair of shoes on the same date of service
- New equipment when current equipment can be modified to accommodate growth
- Nutritional formula representing only a liquid form of food
- Nutritional puddings/bars
- Over-the-counter shoe inserts
- Peri-wash

- Portable oxygen, when oxygen is ordered to be used at night only
- Power tilt-in-space or reclining wheelchairs for a longterm care resident because there is limited staffing
- Pressure gradient garments for maternity-related edema
- Prosthetic appliances for a beneficiary with a potential functional level of K0
- Regular or dietetic foods (e.g., Slimfast, Carnation instant breakfast, etc.)
- Room dehumidifiers
- School Items (e.g., computers, writing aids, book holder, mouse emulator, etc.)
- Second units for school use
- Second wheelchair for beneficiary preference or convenience
- Sensory Devices (e.g., games, toys, etc.)
- Sports drinks/juices
- Stair lifts
- Standard infant/toddler formula
- Therapy modalities (bolsters, physio-rolls, therapy balls, jett mobile)
- Thickeners for foods or liquids (e.g., Thick it)
- Toothettes
- Transcutaneous Nerve Stimulator when prescribed for headaches, visceral abdominal pain, pelvic pain, or temporal mandibular joint (TMJ) pain
- Ultrasonic osteogenesis stimulators
- UV lighting for Seasonal Affective Disorder
- Vacu-brush toothbrushes
- Weight loss or "light" products
- Wheelchair lifts or ramps for home or vehicle (all types)
- Wheelchair accessories (e.g., horns, lights, bags, special colors, etc.)
- Wigs for hair loss

For specific procedure codes that are not covered, refer to the MDCH Medical Supplier Database on the MDCH website or the Coverage Conditions and Requirements Section of this chapter.

(emphasis added by ALJ)

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* * *

The material facts in this case are not in dispute. The Appellant's doctor has requested prior authorization for an ultrasonic osteogenesis stimulator. This is a non-covered item according to the Medicaid Provider Manual policy set forth above. is not required to provide coverage for items excluded by Medicaid Policy under its contract.

The Appellant stated she wants to avoid surgery in order to repair her broken toe and it makes more sense to use the bone stimulator than have to subject herself to surgery.

While this Administrative Law Judge sympathizes with the Appellant's circumstances, this ALJ must uphold the denial issued by contract and Medicaid policy do not require them to cover the requested medical device. This does not mean that the Appellant would not benefit from the medical equipment requested or that she is not deserving of them, but only that the Medicaid policy does not require coverage. The authority possessed by this ALJ does not include equitable determinations nor can policy be disregarded in this forum.

DECISION AND ORDER

The Administrative Law Judge, based on the above findings of fact and conclusions of law, decides that the denial of the Appellant's request for prior-authorization for the ultrasonic osteogenesis stimulator was supported by Medicaid Policy.

IT IS THEREFORE ORDERED that:

The QHP's decision is AFFIRMED.

Jennifer Isiogu Administrative Law Judge for Olga Dazzo, Director Michigan Department of Community Health

CC:



Date Mailed: 4/11/2011

*** NOTICE ***

The State Office of Administrative Hearings and Rules may order a rehearing on either its own motion or at the request of a party within 30 days of the mailing date of this Decision and Order. The State Office of Administrative Hearings and Rules will not order a rehearing on the Department's motion where the final decision or rehearing cannot be implemented within 90 days of the filing of the original request. The Appellant may appeal the Decision and Order to Circuit Court within 30 days of the receipt of the Decision and Order or, if a timely request for rehearing was made, within 30 days of the receipt of the rehearing decision.