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AN ACT relating to utilization reviews.

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2	Be it	t enacted by the General Assembly of the Commonwealth of Kentucky:
3		→ Section 1. KRS 304.17A-611 is amended to read as follows:
4	<u>(1)</u>	A utilization review decision shall not retrospectively deny coverage for health care
5		services provided to a covered person when prior approval has been obtained from
6		the insurer or its designee for those services, unless the approval was based upon
7		fraudulent, materially inaccurate, or misrepresented information submitted by the
8		covered person, authorized person, or the provider.
9	<u>(2)</u>	An insurer shall not require or conduct a prospective or concurrent review for a
0		prescription drug that:
1		(a) Is used in the treatment of alcohol or opioid use disorder; and
2		(b) Contains Methadone, Buprenorphine, or Naltrexone.
3		→ Section 2. KRS 205.536 is amended to read as follows:
4	(1)	A Medicaid managed care organization shall have a utilization review plan, as
5		defined in KRS 304.17A-600, that meets the requirements established in 42 C.F.R.
6		pts. 431, 438, and 456. If the Medicaid managed care organization utilizes a private
7		review agent, as defined in KRS 304.17A-600, the agent shall comply with all
8		applicable requirements of KRS 304.17A-600 to 304.17A-633.
9	(2)	In conducting utilization reviews for Medicaid benefits, each Medicaid managed
20		care organization shall use the medical necessity criteria selected by the Department
21		of Insurance pursuant to KRS 304.38-240, for making determinations of medical
22		necessity and clinical appropriateness pursuant to the utilization review plan
23		required by subsection (1) of this section.
24	<u>(3)</u>	The Department for Medicaid Services or any managed care organization
25		contracted to provide Medicaid benefits pursuant to KRS Chapter 205 shall not
26		require or conduct a prospective or concurrent review, as defined in KRS
27		304.17A-600, for a prescription drug that:

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- 1 (a) Is used in the treatment of alcohol or opioid use disorder; and
- 2 (b) Contains Methadone, Buprenorphine, or Naltrexone.
- 3 → Section 3. This Act takes effect January 1, 2021.