April 15, 2020

Melody Anthony
Medicaid Director
Oklahoma Health Care Authority
Federal Authorities Unit
4345 N. Lincoln Blvd.
Oklahoma City, Oklahoma 73105

Re: SoonerCare 2.0 Section 1115 Demonstration Application

Dear Director Anthony:

The Epilepsy Foundation and Epilepsy Foundation Oklahoma appreciate the opportunity to submit comments on the SoonerCare 2.0 Section 1115 Demonstration Application.

The Epilepsy Foundation is the leading national voluntary health organization that speaks on behalf of more than three million Americans with epilepsy and seizures. Our local chapter, Epilepsy Foundation Oklahoma, provides services and advocates for the approximately 41,100 Oklahomans living with epilepsy and seizures. Together, we foster the wellbeing of children and adults affected by seizures through research programs, educational activities, advocacy, and direct services. Epilepsy is a medical condition that produces seizures, which are sudden electrical surges in the brain, affecting a variety of mental and physical functions. Approximately 1 in 26 Americans will develop epilepsy at some point in their lifetime. For the majority of people living with epilepsy, epilepsy medications are the most common and most cost effective treatment for controlling and/or reducing seizures.

The purpose of the Medicaid program is to provide healthcare coverage for low-income individuals and families, and Epilepsy Foundation and Epilepsy Foundation Oklahoma are committed to ensuring that SoonerCare provides quality and affordable healthcare coverage. We strongly support the State Plan Amendment (SPA) submitted on March 6, 2020 that will expand Medicaid coverage in Oklahoma to individuals making less than 138 percent of the federal poverty level ($2,498 for a family of three) beginning July 1, 2020. We urge the state to implement this expansion as quickly as possible.

Unfortunately the Soonercare 2.0 proposal would greatly reduce the benefits of Oklahoma’s Medicaid expansion and jeopardize beneficiaries’ access to care. This proposal would shift the program to a capped funding structure, reduce patients’ access to critical benefits and services, and add new administrative and financial barriers to the program that would undoubtedly lead to coverage losses – the state itself estimates 7,600 fewer individuals would be enrolled due to the premium and work requirements in the first year of the demonstration alone. These coverage losses are unacceptable to Epilepsy Foundation and Epilepsy Foundation Oklahoma.
Epilepsy Foundation and Epilepsy Foundation Oklahoma would oppose this proposal under any circumstances, but it is especially dangerous to move forward with this proposal during a public health emergency such as the current COVID-19 pandemic. Now, more than ever, it is critical to ensure that individuals living with epilepsy and seizures have meaningful access to care and treatment options in order to control their seizures. People with epilepsy who suddenly lose access to their treatment options are at a greater risk of accident, injury, and hospitalization, further taxing an already stretched health care resources in the state. Epilepsy Foundation and Epilepsy Foundation Oklahoma urge the state to immediately withdraw this proposal.

**Block Grant Structure**

While the state uses an application template for its proposal which is to be used by states “applying to use either an aggregate or a per capita cap financing model for certain populations” the proposal includes no details about the cap, how it would work or how much capped funding the state would receive. Epilepsy Foundation and Epilepsy Foundation Oklahoma are extremely concerned with the lack of detail in Oklahoma’s proposal. Such a drastic change in Oklahoma’s Medicaid program will undoubtedly have a dramatic impact on patients, but without additional details, it is difficult for Epilepsy Foundation to fully comment on all of the possible impacts of a block grant or per capita cap on the patients we represent.

We oppose the use of block grants and per capita caps in the Medicaid program. Neither financing structure will protect either the state or patients from enormous financial risk. As the gap between the capped allotment and actual costs of patient care increases over time, states will likely limit enrollment, reduce benefits, lower provider payments or increase cost-sharing, all of which would cause significant harm to the patients we represent.

Many situations could lead Oklahoma to exceed a funding cap. A public health emergency like COVID-19 will greatly increase healthcare costs above negotiated caps and an economic recession would similarly increase enrollment in and costs associated with SoonerCare. Allowing states to renegotiate their caps due to special circumstances like a public health emergency will not protect states and patients from the financial risks of block grants; there is no guarantee that CMS would authorize additional funding via a potentially lengthy re-negotiation process and, in the interim, Oklahoma would face strong incentives to make cuts to the Medicaid program that would harm patients. Moreover, the federal government will likely be focused on responding to the emergency at hand – putting the renegotiation of complex budget neutrality agreements on the back burner. Again, while it would be dangerous for Oklahoma to pursue a capped funding structure at any time, it is especially dangerous to do so now.

**Barriers to Coverage**

**Retroactive Coverage**

Oklahoma has requested the authority to waive retroactive eligibility, a policy that prevents gaps in coverage by covering individuals for up to 90 days prior to the month of application, assuming the individual is eligible for
Medicaid coverage during that timeframe. It is common that individuals are unaware they are eligible for Medicaid until a medical event or diagnosis occurs. Retroactive eligibility allows patients who have been diagnosed with a serious illness, such as epilepsy, to begin treatment without being burdened by medical debt prior to their official eligibility determination.

Medicaid paperwork can be burdensome and often times confusing. A Medicaid enrollee may not have understood or received a notice of Medicaid renewal and only discovered the coverage lapse when picking up a prescription or going to see their doctor. Without retroactive eligibility, Medicaid enrollees could then face substantial costs at their doctor’s office or pharmacy. When Ohio was considering a similar provision in 2016, one estimate predicted that hospitals could accrue as much as $2.5 billion more in uncompensated care as a result of the waiver. Additional uncompensated care would be especially problematic at the current time because it would add to the financial challenges hospitals are facing as a result of COVID-19. Epilepsy Foundation and Epilepsy Foundation Oklahoma oppose a waiver of retroactive coverage.

**Premiums and Cost-Sharing**

Under Oklahoma’s application, individuals with incomes above 42 percent of the federal poverty level would have to pay premiums ranging from $5 to $15 per month. Individuals could not enroll in coverage until they pay their first premium and could lose their coverage if they are unable to pay future premiums. This policy would likely both increase the number of enrollees who lose Medicaid coverage and also discourage eligible people from enrolling in the program. For example, when Oregon implemented a premium in its Medicaid program, with a maximum premium of $20 per month, almost half of enrollees lost coverage. Epilepsy Foundation and Epilepsy Foundation Oklahoma believe that these premiums will create significant financial barriers for patients that jeopardize their access to needed care and therefore opposes this policy.

Oklahoma’s application also includes copays for its Medicaid program, including an $8 copay for non-emergent use of the emergency department (ED). This policy could deter people from seeking necessary care during an emergency. Furthermore, evidence suggests this type of cost sharing may not result in the intended cost savings. For example, a study of enrollees in Oregon’s Medicaid program demonstrated that implementation of a copay on emergency services resulted in decreased utilization of such services but did not result in cost savings because of subsequent use of more intensive and expensive services. Epilepsy Foundation and Epilepsy Foundation Oklahoma oppose this punitive proposal for a $8 copayment for non-emergent use of the ED.

Finally, Oklahoma requests the authority to increase premiums and cost-sharing up to five percent of household income. This would put an enormous financial burden on patients that would again jeopardize their coverage. Additionally, any future increases in cost-sharing should go through a full public comment process and review by CMS, which are important opportunities for the public to provide feedback on how the program is working for key stakeholders before any policies are implemented or continued. It is especially important that beneficiaries impacted by the demonstration waiver have the ability to provide feedback to the state and CMS. Epilepsy Foundation and Epilepsy Foundation Oklahoma urge Oklahoma to withdraw this request.
**Work Requirements**

Under the application, individuals between the ages of 19 and 60 be required to prove that they work up to 80 hours per month or meet exemptions. One major consequence of this proposal will be to increase the administrative burden on individuals in the Medicaid program. Increasing administrative requirements will likely decrease the number of individuals with Medicaid coverage, regardless of whether they are exempt or not. For example, when Arkansas implemented a similar policy, the state terminated coverage for over 18,000 individuals, and in New Hampshire, nearly 17,000 individuals would have lost coverage if the state had not suspended implementation of its requirement. The U.S. Court of Appeals for the District of Columbia recently reaffirmed that the purpose of the Medicaid program is to provide healthcare coverage and that Arkansas’ restrictive waiver, including the work requirement policy, did not meet that objective.

Failing to navigate these burdensome administrative requirements could have serious – even life or death – consequences for people with serious, acute and chronic diseases. If the state finds that individuals have failed to comply with the new requirements after one month, they will be disenrolled from coverage.

Epilepsy Foundation and Epilepsy Foundation Oklahoma are also concerned that the current exemption criteria may not capture all individuals with, or at risk of, serious and chronic health conditions that prevent them from working. Regardless, even exempt enrollees may have to report their exemption, creating opportunities for administrative error that could jeopardize their coverage. In Arkansas, many individuals were unaware of the new requirements and therefore unaware that they needed to apply for such an exemption. No exemption criteria can circumvent this problem and the serious risk to the health of the people we represent.

Administering these requirements will also be expensive for the state of Oklahoma. States such as Kentucky, Tennessee and Virginia have estimated that setting up the administrative systems to track and verify exemptions and work activities will cost tens of millions of dollars. This would divert federal resources from Medicaid’s core goal – providing health coverage to those without access to care – and compromise the fiscal health of Indiana’s Medicaid program.

The evidence is clear that most people on Medicaid who can work already do so, and those who are unable to work often have physical or mental health conditions that interfere with their ability to work. Evaluations of Arkansas’s waiver demonstrate that it did not lead to increased employment among the Medicaid population. In contrast, continuous Medicaid coverage can actually help people find and sustain employment. For example, a report examining Medicaid expansion in Ohio found that the majority of enrollees reported that being enrolled in Medicaid made it easier to work or look for work (83.5 percent and 60 percent, respectively). Terminating individuals’ Medicaid coverage for non-compliance with work requirements will hurt rather than help people search for and obtain employment.
Benefit Package

Oklahoma’s application also jeopardizes access to vital services for low-income patients served by the Medicaid program, particularly those with serious and chronic diseases.

Oklahoma’s application proposes to waive Early and Periodic Screening, Diagnosis and Treatment (EPSDT) for individuals aged 19 and 20. EPSDT provides access to critical services and treatments for kids and young adults living in poverty. As these young adults transition to higher education or jobs, it is important that they receive the same medical care for any illness or chronic disease they might have. Disruption in medical treatment could have negative consequences for their long-term health and economic security. Epilepsy Foundation and Epilepsy Foundation Oklahoma oppose this provision.

Oklahoma has also requested to eliminate Non-Emergency Medical Transportation (NEMT) benefits. Low-income patients may not own a car and may lack access to reliable public transportation, especially in rural areas. Removing this benefit will therefore harm patients who need to attend regular visits with their providers to manage their medications and treatments. For example, one study found patients with asthma, hypertension or heart disease who needed multiple visits to a medical professional more likely to keep their appointments if they had NEMT.

Epilepsy Foundation and Epilepsy Foundation Oklahoma strongly oppose this policy.

Finally, the proposal also states that Oklahoma will “continue to investigate the potential benefits of a limited prescription drug formulary and request the flexibility to make changes to our prescription drug benefit, following appropriate advance notice procedures.” We believe that any changes to the prescription drug benefit that limit access to medications will be detrimental to individuals living with epilepsy and seizures. Prescription drugs have different indications, different mechanisms of action, and different side effects, depending on the person’s diagnosis and comorbidities. Treatment for epilepsy is not “one size fits all,” and individuals must have access to the broad range of therapeutic options in order to find the right treatment for them in order to achieve zero seizures with zero side effects. Restricting prescription drug benefits would limit the ability of providers to make the best medical decisions for the care of their patients.

Public Comment Process

A robust public comment process is an essential component on any Section 1115 demonstration proposal. Unfortunately, Oklahoma’s public comment process has been rushed and it has been difficult for individuals to participate. This has made meaningful comment impossible for many critical stakeholders.

The core objective of the Medicaid program is to furnish healthcare to low-income populations. This demonstration application does not further that goal and Epilepsy Foundation and Epilepsy Foundation Oklahoma oppose the proposal. We instead urge Oklahoma to move forward with implementing its SPA for Medicaid expansion as soon as possible without imposing any additional barriers to coverage. Thank you for the opportunity to submit comments.
Sincerely,

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Acting State Director  
Epilepsy Foundation Oklahoma

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Vice President, Government Relations & Advocacy  
Epilepsy Foundation

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