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## Fast-Track Regulation Agency Background Document

<b>Agency name</b>	Department of Medical Assistance Services
<b>Virginia Administrative Code (VAC) citation(s)</b>	12 VAC 30-80-30
<b>Regulation title(s)</b>	Methods and Standards for Establishing Payment Rate; Other Types of Care
<b>Action title</b>	Supplemental Payments for Type One Physicians
<b>Date this document prepared</b>	October 30, 2017

This information is required for executive branch review and the Virginia Registrar of Regulations, pursuant to the Virginia Administrative Process Act (APA), Executive Orders 17 (2014) and 58 (1999), and the *Virginia Register Form, Style, and Procedure Manual*.

### Brief summary

*Please provide a brief summary (preferably no more than 2 or 3 paragraphs) of the proposed new regulation, proposed amendments to the existing regulation, or the regulation proposed to be repealed. Alert the reader to all substantive matters or changes. If applicable, generally describe the existing regulation.*

12-VAC 30-80-30(16) identifies the requirements for supplemental payments provided to Type One physicians. This regulatory action updates the supplemental payment amounts for Type One physicians effective April 1, 2017 and May 1, 2017.

**Statement of final agency action**

Please provide a statement of the final action taken by the agency including: 1) the date the action was taken; 2) the name of the agency taking the action; and 3) the title of the regulation.

I hereby approve the foregoing Regulatory Review Summary entitled Supplemental Payments for Type One Physicians with the attached amended regulations (12 VAC 30-80-30: Methods and Standards for Establishing Payment Rate; Other Types of Care) and adopt the action stated therein. I certify that this fast track regulatory action has completed all the requirements of the Code of Virginia § 2.2-4012.1, of the Administrative Process Act.

10-30-2017

/Cynthia B. Jones/

Date

Cynthia B. Jones, Director

Dept. of Medical Assistance Services

**Legal basis**

Please identify the state and/or federal legal authority to promulgate this proposed regulation, including: 1) the most relevant citations to the Code of Virginia or General Assembly chapter number(s), if applicable; and 2) promulgating entity, i.e., agency, board, or person. Your citation should include a specific provision authorizing the promulgating entity to regulate this specific subject or program, as well as a reference to the agency/board/person’s overall regulatory authority.

The *Code of Virginia* (1950) as amended, § 32.1-325, grants to the Board of Medical Assistance Services the authority to administer and amend the Plan for Medical Assistance. The *Code of Virginia* (1950) as amended, § 32.1-324, authorizes the Director of DMAS to administer and amend the Plan for Medical Assistance according to the Board's requirements. The Medicaid authority as established by § 1902 (a) of the *Social Security Act* [42 U.S.C. 1396a] provides governing authority for payments for services.

In addition, authority for this change is provided in the 2017 *Acts of Assembly, Chapter 836, Item 306.B.4* which states:

*“The Department of Medical Assistance Service shall have the authority to increase Medicaid payments for Type One hospitals and physicians consistent with the appropriations to compensate for limits on disproportionate share hospital (DSH) payments to Type One hospitals that the department would otherwise make.”*

### Purpose

*Please explain the need for the new or amended regulation. Describe the rationale or justification of the proposed regulatory action. **Describe the specific reasons the regulation is essential to protect the health, safety or welfare of citizens.** Discuss the goals of the proposal and the problems the proposal is intended to solve.*

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The purpose of this action is to update the physician supplemental payments for Type One physicians. The supplemental payment calculation amount in the current regulatory text was effective April 8, 2014. This change allows for updated supplemental payments to Type One physicians which are expected to improve access to services for Medicaid recipients.

### Rationale for using fast-track process

*Please **explain the rationale for using the fast-track process** in promulgating this regulation. Why do you expect this rulemaking to be noncontroversial?*

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This regulatory action is being promulgated as a fast track action because it is not expected to be controversial. The fiscal or budgetary impacts to DMAS are already provided in the agency’s appropriations. These changes have been mandated by the Appropriations Act and CMS.

### Substance

*Please briefly identify and explain the new substantive provisions, the substantive changes to existing sections, or both. A more detailed discussion is provided in the “Detail of changes” section below.*

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This regulatory change is intended to update the physician supplemental payments for physician practice plans affiliated with Type One hospitals (state academic health systems). A Type One physician is a member of a practice group organized by or under the control of a state academic health system or an academic health system that operates under a state authority and includes a hospital, which has entered into contractual agreements for the assignment of payments in accordance with 42 CFR 447.10. These payments are calculated as the difference between the maximum payment allowed and regular payments. The Center for Medicare and Medicaid Services (CMS) has determined that the maximum allowed is the average commercial rate (ACR).

This action will update the maximum rate to 256% of the Medicare rate effective April 1, 2017, and 258% effective May 1, 2017 based on the most recent information on the ACR furnished by the state academic health systems and consistent with appropriate prior public notices.

### Issues

*Please identify the issues associated with the proposed regulatory action, including: 1) the primary advantages and disadvantages to the public, such as individual private citizens or businesses, of implementing the new or amended provisions; 2) the primary advantages and disadvantages to the agency or the Commonwealth; and 3) other pertinent matters of interest to the regulated community, government officials, and the public. If there are no disadvantages to the public or the Commonwealth, please indicate.*

These changes create no disadvantages to the public, the Agency, the Commonwealth, or the regulated community. The changes all implement directives in the state budget and update existing regulations to conform with the State Plan.

Updating supplemental payment amounts for Type One physicians is expected to be advantageous as it will improve access to services.

### Requirements more restrictive than federal

*Please identify and describe any requirement of the proposal which is more restrictive than applicable federal requirements. Include a rationale for the need for the more restrictive requirements. If there are no applicable federal requirements or no requirements that exceed applicable federal requirements, include a statement to that effect.*

There are no requirements more restrictive than federal contained in these recommendations.

### Localities particularly affected

*Please identify any locality particularly affected by the proposed regulation. Locality particularly affected means any locality which bears any identified disproportionate material impact which would not be experienced by other localities.*

There will be no localities that are more affected than others as these requirements will apply statewide.

### Regulatory flexibility analysis

*Pursuant to § 2.2-4007.1B of the Code of Virginia, please describe the agency's analysis of alternative regulatory methods, consistent with health, safety, environmental, and economic welfare, that will accomplish the objectives of applicable law while minimizing the adverse impact on small business. Alternative regulatory methods include, at a minimum: 1) the establishment of less stringent compliance or reporting requirements; 2) the establishment of less stringent schedules or deadlines for compliance or reporting requirements; 3) the consolidation or simplification of compliance or reporting requirements; 4) the establishment of performance standards for small businesses to replace design or operational standards required in the proposed regulation; and 5) the exemption of small businesses from all or any part of the requirements contained in the proposed regulation.*

This regulatory action is not expected to affect small businesses as it does not impose compliance or reporting requirements, nor deadlines for reporting, nor does it establish performance standards to replace design or operational standards.

**Economic impact**

*Please identify the anticipated economic impact of the proposed new regulations or amendments to the existing regulation. When describing a particular economic impact, please specify which new requirement or change in requirement creates the anticipated economic impact.*

<b>Projected cost to the state to implement and enforce the proposed regulation, including:</b> a) fund source / fund detail; and b) a delineation of one-time versus on-going expenditures	No net additional cost as DMAS obligations can be reduced in other areas.
<b>Projected cost of the new regulations or changes to existing regulations on localities.</b>	None
<b>Description of the individuals, businesses, or other entities likely to be affected by the new regulations or changes to existing regulations.</b>	Physicians in practice plans affiliated with Type One hospitals.
<b>Agency’s best estimate of the number of such entities that will be affected. Please include an estimate of the number of small businesses affected. Small business means a business entity, including its affiliates, that:</b> a) is independently owned and operated and; b) employs fewer than 500 full-time employees or has gross annual sales of less than \$6 million.	Two practice plans employing 350 physicians.
<b>All projected costs of the new regulations or changes to existing regulations for affected individuals, businesses, or other entities. Please be specific and include all costs including:</b> a) the projected reporting, recordkeeping, and other administrative costs required for compliance by small businesses; and b) specify any costs related to the development of real estate for commercial or residential purposes that are a consequence of the proposed regulatory changes or new regulations.	None
<b>Beneficial impact the regulation is designed to produce.</b>	Access to physicians in practice plans affiliated with Type One hospitals.

### Alternatives

*Please describe any viable alternatives to the proposal considered and the rationale used by the agency to select the least burdensome or intrusive alternative that meets the essential purpose of the action. Also, include discussion of less intrusive or less costly alternatives for small businesses, as defined in § 2.2-4007.1 of the Code of Virginia, of achieving the purpose of the regulation.*

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No other alternative would meet the requirements set forth in the 2017 Acts of Assembly.

### Public participation notice

*If an objection to the use of the fast-track process is received within the 30-day public comment period from 10 or more persons, any member of the applicable standing committee of either house of the General Assembly or of the Joint Commission on Administrative Rules, the agency shall: 1) file notice of the objections with the Registrar of Regulations for publication in the Virginia Register; and 2) proceed with the normal promulgation process with the initial publication of the fast-track regulation serving as the Notice of Intended Regulatory Action.*

### Family Impact

*Please assess the impact of the proposed regulatory action on the institution of the family and family stability including to what extent the regulatory action will: 1) strengthen or erode the authority and rights of parents in the education, nurturing, and supervision of their children; 2) encourage or discourage economic self-sufficiency, self-pride, and the assumption of responsibility for oneself, one's spouse, and one's children and/or elderly parents; 3) strengthen or erode the marital commitment; and 4) increase or decrease disposable family income.*

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These changes do not strengthen or erode the authority or rights of parents in the education, nurturing, and supervision of their children; nor encourage or discourage economic self-sufficiency, self-pride, and the assumption of responsibility for oneself, one's spouse, and one's children and/or elderly parents. It does not strengthen or erode the marital commitment, but may decrease disposable family income depending upon which provider the recipient chooses for the item or service prescribed.

### Detail of changes

*Please list all changes that are being proposed and the consequences of the proposed changes; explain the new requirements and what they mean rather than merely quoting the proposed text of the regulation. If the proposed regulation is a new chapter, describe the intent of the language and the expected impact. Please describe the difference between existing regulation(s) and/or agency practice(s) and what is being proposed in this regulatory action. If the proposed regulation is intended to replace an emergency regulation, please list separately: (1) all differences between the **pre-emergency** regulation and this proposed regulation; and 2) only changes made since the publication of the emergency regulation.*

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Current section number	Proposed new section number, if applicable	Current requirement	Proposed change, intent, rationale, and likely impact of proposed requirements
12 VAC 30-80-30 (16)		The current supplemental payment amount for Type One physician services which was effective April 8, 2014 is the difference between the Medicaid payments otherwise made for physician services and 201% of Medicare rates.	The proposed change updates the current supplemental payment amount for Type One physician services to the difference between the Medicaid payments otherwise made for physician services and 256% of Medicare rates effective April 1, 2017. Effective May 1, 2017, the supplemental payment amount increases to the difference between the Medicaid payments otherwise made for physician services and 258% of Medicare rates.