

authority and consistent with the law to further achieve these aims.

II. Solicitation of Comments

HHS is interested in soliciting public comments about changes to existing regulations or guidance, or other actions within HHS's authority, that could further the following goals with respect to the individual and small group health insurance markets:

1. *Empowering patients and promoting consumer choice.* What activities would best inform consumers and help them choose a plan that best meets their needs? Which regulations currently reduce consumer choices of how to finance their health care and health insurance needs? Choice includes the freedom to choose how to finance one's healthcare, which insurer to use, and which provider to use.

2. *Stabilizing the individual, small group, and non-traditional health insurance markets.* What changes would bring stability to the risk pool, promote continuous coverage, increase the number of younger and healthier consumers purchasing plans, reduce uncertainty and volatility, and encourage uninsured individuals to buy coverage?

3. *Enhancing affordability.* What steps can HHS take to enhance the affordability of coverage for individual consumers and small businesses?

4. *Affirming the traditional regulatory authority of the States in regulating the business of health insurance.* Which HHS regulations or policies have impeded or unnecessarily interfered with States' primary role in regulating the health insurance markets they know best?

This is a request for information only. Respondents are encouraged to provide complete but concise responses to the questions outlined above. We note that a response to every question is not required. This request for information is issued solely for information and planning purposes; it does not constitute a notice of proposed rulemaking or request for proposals, applications, proposal abstracts, or quotations. This request for information does not commit the United States Government ("Government") to contract for any supplies or services or make a grant award. Further, HHS is not seeking proposals through this request for information and will not accept unsolicited proposals. Respondents are advised that the Government will not pay for any information or administrative costs incurred in response to this request for information; all costs associated with responding to this request for information will be

solely at the interested party's expense. Not responding to this request for information does not preclude participation in any future rulemaking or procurement, if conducted. It is the responsibility of the potential responders to monitor this request for information announcement for additional information pertaining to this request. We also note that HHS will not respond to questions about the policy issues raised in this request for information. HHS may or may not choose to contact individual responders. Such communications would only serve to further clarify written responses. Contractor support personnel may be used to review request for information responses. Responses to this notice are not offers and cannot be accepted by the Government to form a binding contract or issue a grant. Information obtained as a result of this request for information may be used by the Government for program planning on a non-attribution basis. Respondents should not include any information that might be considered proprietary or confidential. This request for information should not be construed as a commitment or authorization to incur cost for which reimbursement would be required or sought. All submissions become Government property and will not be returned. HHS may publically post the comments received, or a summary thereof. While responses to this request for information do not bind HHS to any further actions related to the response, all submissions will be made publicly available on <http://www.regulations.gov>.

III. Collection of Information Requirements

This document does not impose information collection requirements, that is, reporting, recordkeeping or third-party disclosure requirements. This request for information constitutes a general solicitation of comments. In accordance with the implementing regulations of the Paperwork Reduction Act (PRA) at 5 CFR 1320.3(h)(4), information subject to the PRA does not generally include "facts or opinions submitted in response to general solicitations of comments from the public, published in the **Federal Register** or other publications, regardless of the form or format thereof, provided that no person is required to supply specific information pertaining to the commenter, other than that necessary for self-identification, as a condition of the agency's full consideration of the comment." Consequently, this document need not be reviewed by the Office of Management and Budget under the

authority of the Paperwork Reduction Act of 1995 (44 U.S.C. 3501 *et seq.*).

Dated: June 6, 2017.

Seema Verma,

Administrator, Centers for Medicare & Medicaid Services.

Dated: June 7, 2017.

Thomas E. Price,

Secretary, Department of Health and Human Services.

[FR Doc. 2017-12130 Filed 6-8-17; 4:15 pm]

BILLING CODE 4120-01-P

FEDERAL COMMUNICATIONS COMMISSION

47 CFR Part 73

[MB Docket No. 11-54; RM-11624; DA 17-510]

Television Broadcasting Services; Augusta, Georgia

AGENCY: Federal Communications Commission.

ACTION: Proposed rule; withdrawal.

SUMMARY: The Commission has before it a petition for rulemaking filed by Southern Media Holdings, Inc. (SMH), the former licensee of WFXG, Augusta, Georgia, requesting the substitution of channel 51 for channel 31 at Augusta. WFXG License Subsidiary, LLC (Licensee) is now the licensee of WFXG. Station WFXG was allotted channel 51 as its post-transition DTV channel and operated a licensed facility on that channel. In 2008, SMH filed a petition for rulemaking requesting that channel 31 be substituted for channel 51, and the Commission granted that request. SMH subsequently requested that the Commission change its channel back to channel 51 and we issued a Notice of Proposed Rulemaking, which was contested. On April 28, 2017, Licensee filed a letter withdrawing its pending request to substitute channel 51 for channel 31, explaining that it had licensed the channel 31 facility and that WFXG was reassigned to channel 36 in connection with the post-incentive auction repacking of the broadcast television spectrum.

DATES: The proposed rule published on April 4, 2011 (76 FR 18497) is withdrawn as of June 12, 2017.

FOR FURTHER INFORMATION CONTACT: Joyce Bernstein, Joyce.Bernstein@fcc.gov, Media Bureau, (202) 418-1647.

SUPPLEMENTARY INFORMATION: This is a synopsis of the Commission's *Order*, MB Docket No. 11-54, adopted May 25, 2017, and released May 25, 2017. The full text of this document is available for

public inspection and copying during normal business hours in the FCC's Reference Information Center at Portals II, CY-A257, 445 12th Street SW., Washington, DC, 20554. This document will also be available via ECFS (<http://fjallfoss.fcc.gov/ecfs/>). To request materials in accessible formats for people with disabilities (braille, large print, electronic files, audio format), send an email to fcc504@fcc.gov or call the Consumer & Governmental Affairs Bureau at 202-418-0530 (voice), 202-418-0432 (tty).

This document does not contain information collection requirements subject to the Paperwork Reduction Act of 1995, Public Law 104-13. In addition, therefore, it does not contain any information collection burden "for small business concerns with fewer than 25 employees," pursuant to the Small Business Paperwork Relief Act of 2002, Public Law 107-198, *see* 44 U.S.C. 3506(c)(4). Provisions of the Regulatory Flexibility Act of 1980, *see* 5 U.S.C. 601-612, do not apply to this proceeding.

The Commission will send a copy of this *Report and Order* in a report to be sent to Congress and the Government Accountability Office pursuant to the Congressional Review Act, *see* 5 U.S.C. 801(a)(1)(A).

List of Subjects in 47 CFR Part 73

Television.

Federal Communications Commission.

Barbara A. Kreisman,

Chief, Video Division, Media Bureau.

[FR Doc. 2017-11947 Filed 6-9-17; 8:45 am]

BILLING CODE 6712-01-P

DEPARTMENT OF TRANSPORTATION

Federal Motor Carrier Safety Administration

49 CFR Part 383

[Docket No. FMCSA-2016-0346]

RIN 2126-AB98

Commercial Learner's Permit Validity

AGENCY: Federal Motor Carrier Safety Administration (FMCSA), DOT.

ACTION: Notice of Proposed Rulemaking (NPRM), request for comments.

SUMMARY: FMCSA proposes to amend the Federal Motor Carrier Safety Regulations (FMCSRs) to allow States to issue a commercial learner's permit (CLP) with an expiration date of up to one year from the date of initial issuance. CLPs issued for shorter periods may be renewed but the total

period of time between the date of initial issuance and the expiration of the renewed CLP could not exceed one year. This proposed amendment would replace the current regulations, which require the States to issue CLPs initially for no more than 180 days, with the possibility of an additional 180-day renewal at the State's discretion.

DATES: Comments on this notice must be received on or before August 11, 2017.

ADDRESSES: You may submit comments identified by Docket Number FMCSA-2016-0346 using any of the following methods:

- *Federal eRulemaking Portal:* <http://www.regulations.gov>. Follow the online instructions for submitting comments.
- *Mail:* Docket Management Facility, U.S. Department of Transportation, 1200 New Jersey Avenue SE., West Building, Ground Floor, Room W12-140, Washington, DC 20590-0001.
- *Hand Delivery or Courier:* West Building, Ground Floor, Room W12-140, 1200 New Jersey Avenue SE., Washington, DC, between 9 a.m. and 5 p.m., Monday through Friday, except Federal holidays.
- *Fax:* 202-493-2251.

To avoid duplication, please use only one of these four methods. See the "Public Participation and Request for Comments" portion of the **SUPPLEMENTARY INFORMATION** section for instructions on submitting comments, including collection of information comments for the Office of Information and Regulatory Affairs, OMB.

FOR FURTHER INFORMATION CONTACT: Mr. Selden Fritschner, CDL Division, Federal Motor Carrier Safety Administration, 1200 New Jersey Avenue SE., Washington, DC 20590-0001, by email at selden.fritschner@dot.gov, or by telephone at 202-366-0677.

SUPPLEMENTARY INFORMATION:

I. Public Participation and Request for Comments

A. Submitting Comments

If you submit a comment, please include the docket number for this NPRM (Docket No. FMCSA-2016-0346), indicate the specific section of this document to which each section applies, and provide a reason for each suggestion or recommendation. You may submit your comments and material online or by fax, mail, or hand delivery but please use only one of these means. FMCSA recommends that you include your name and a mailing address, an email address, or a phone number in the body of your document so that FMCSA can contact you if there

are questions regarding your submission.

To submit your comment online, go to <http://www.regulations.gov>, put the docket number, FMCSA-2016-0346, in the keyword box, and click "Search." When the new screen appears, click on the "Comment Now!" button and type your comment into the text box on the following screen. Choose whether you are submitting your comment as an individual or on behalf of a third party and then submit.

If you submit your comments by mail or hand delivery, submit them in an unbound format, no larger than 8½ by 11 inches, suitable for copying and electronic filing. If you submit comments by mail and would like to know that they reached the facility, please enclose a stamped, self-addressed postcard or envelope.

FMCSA will consider all comments and material received during the comment period and may change this proposed rule based on your comments. FMCSA may issue a final rule at any time after the close of the comment period.

Confidential Business Information

Confidential Business Information (CBI) is commercial or financial information that is customarily not made available to the general public by the submitter. Under the Freedom of Information Act, CBI is eligible for protection from public disclosure. If you have CBI that is relevant or responsive to this NPRM, it is important that you clearly designate the submitted comments as CBI. Accordingly, please mark each page of your submission as "confidential" or "CBI." Submissions designated as CBI and meeting the definition noted above will not be placed in the public docket of this NPRM. Submissions containing CBI should be sent to Brian Dahlin, Chief, Regulatory Analysis Division, 1200 New Jersey Avenue SE., Washington, DC 20590. Any commentary that FMCSA receives which is not specifically designated as CBI will be placed in the public docket for this rulemaking.

FMCSA will consider all comments and material received during the comment period.

B. Viewing Comments and Documents

To view comments, as well as any documents mentioned in this preamble as being available in the docket, go to <http://www.regulations.gov>. Insert the docket number, FMCSA-2016-0346, in the keyword box, and click "Search." Next, click the "Open Docket Folder" button and choose the document to review. If you do not have access to the