§ 94.29 Restrictions on importation of fresh (chilled or frozen) beef and ovine meat from specified regions.

Notwithstanding any other provisions of this part, fresh (chilled or frozen) beef from a region in Argentina located north of Patagonia South and Patagonia North B, referred to as Northern Argentina, and fresh (chilled or frozen) beef and ovine meat from Uruguay may be exported to the United States under the following conditions:

(a) The meat is beef or ovine meat from animals that have been born, raised, and slaughtered in the exporting region of Argentina or in Uruguay.
(b) Foot-and-mouth disease has not been diagnosed in the exporting region of Argentina or in Uruguay.
(c) The meat comes from bovines or sheep that originated from premises where foot-and-mouth disease has not been present during the lifetime of any bovines and sheep slaughtered for the export of beef and ovine meat to the United States.
(d) The meat comes from bovines or sheep that were moved directly from the premises of origin to the slaughtering establishment without any contact with other animals.
(e) The meat comes from bovines or sheep that received ante-mortem and post-mortem veterinary inspections, paying particular attention to the head and feet, at the slaughtering establishment, with no evidence found of vesicular disease.
(f) The meat consists only of bovine parts or ovine parts that are, by standard practice, part of the animal’s carcass that is placed in a chiller for maturation after slaughter. The bovine and ovine parts that may not be imported include all parts of the head, feet, hump, hooves, and internal organs.
(g) All bone and visually identifiable blood clots and lymphoid tissue have been removed from the meat.
(h) The meat has not been in contact with meat from regions other than those listed in § 94.1(a).
(i) The meat came from bovine carcasses that were allowed to mature at 40 to 50 °F (4 to 10 °C) for a minimum of 24 hours after slaughter and that reached a pH below 6.0 in the loin muscle at the end of the maturation period. Measurements for pH must be taken at the middle of both longissimus dorsi muscles. Any carcass in which the pH does not reach less than 6.0 may be allowed to mature an additional 24 hours and be retested, and, if the carcass still has not reached a pH of less than 6.0 after 48 hours, the meat from the carcass may not be exported to the United States.
(j) An authorized veterinary official of the government of the exporting region certifies on the foreign meat inspection certificate that the above conditions have been met.
(k) The establishment in which the bovines and sheep are slaughtered allows periodic on-site evaluation and subsequent inspection of its facilities, records, and operations by an APHIS representative.

(Authorized by the Office of Management and Budget under control number 0579–0372)

Done in Washington, DC, this 26th day of August 2014.

Michael C. Greigre, Acting Administrator, Animal and Plant Health Inspection Service.

[FR Doc. 2014–20643 Filed 8–28–14; 8:45 am]

BILLING CODE 3410–34–P

FEDERAL TRADE COMMISSION

16 CFR Part 312

RIN 3084–AB20

Children’s Online Privacy Protection Rule: AgeCheq Application for Parental Consent Method

AGENCY: Federal Trade Commission (FTC or Commission).

ACTION: Request for public comment.

SUMMARY: The Federal Trade Commission publishes this request for public comment concerning the proposed parental consent method submitted by AgeCheq Inc. (“AgeCheq”) under the Voluntary Commission Approval Processes provision of the Children’s Online Privacy Protection Rule.

DATES: Written comments must be received on or before September 30, 2014.

ADDRESS: Interested parties may file a comment online or on paper, by following the instructions in the Request for Comment part of the SUPPLEMENTARY INFORMATION section below. Write “AgeCheq Application for Parental Consent Method, Project No. P–145410” on your comment, and file your comment online at https://ftcpublic comentárioworks.com/ftc/coppaagecheqapp by following the instructions on the web-based form. If you prefer to file your comment on paper, mail your comment to the following address: Federal Trade Commission, Office of the Secretary, Constitution Center, 400 7th Street SW., 5th Floor, Suite 5610 (Annex K), Washington, DC 20024.

FOR FURTHER INFORMATION CONTACT:


SUPPLEMENTARY INFORMATION:

Section A. Background

On October 20, 1999, the Commission issued its final Rule1 pursuant to the Children’s Online Privacy Protection Act, 15 U.S.C. 6501 et seq., which became effective on April 21, 2000.2 On December 19, 2012, the Commission amended the Rule, and these amendments became effective on July 1, 2013.3 The Rule requires certain Web site operators to post privacy policies and provide notice, and to obtain verifiable parental consent, prior to collecting, using, or disclosing personal information from children under the age of 13. The Rule enumerates methods for obtaining verifiable parental consent, while also allowing an interested party to file a written request for Commission approval of parental consent methods not currently enumerated.4 To be considered, the party must submit a detailed description of the proposed parental consent method, together with an analysis of how the method meets the requirements for parental consent described in 16 CFR 312.5(b)(1).

Pursuant to § 312.12(a) of the Rule, AgeCheq has submitted a proposed parental consent method to the Commission for approval. The full text of its application is available on the Commission’s Web site at www.ftc.gov.

Section B. Questions on the Parental Consent Method

The Commission is seeking comment on the proposed parental consent method, and is particularly interested in receiving comment on the questions that follow. These questions are designed to assist the Commission’s consideration of the petition and should not be construed as a limitation on the issues on which public comment may be submitted. Responses to these questions should cite the number of the question being answered. For all comments submitted, please provide any relevant data, statistics, or any other evidence, upon which those comments are based.

1 64 FR 59888 (1999).
2 16 CFR part 312.
3 78 FR 3972 (2013).
4 16 CFR 312.12(a); 78 FR at 3991–3992, 4013.
1. Is this method, both with respect to the process for obtaining consent for an initial operator and any subsequent operators, already covered by existing methods enumerated in § 312.5(b)(1) of the Rule?

2. If this is a new method, provide comments on whether the proposed parental consent method, both with respect to an initial operator and any subsequent operators, meets the requirements for parental consent laid out in 16 CFR 312.5(b)(1). Specifically, the Commission is looking for comments on whether the proposed parental consent method is reasonably calculated, in light of available technology, to ensure that the person providing consent is the child’s parent.

3. Does this proposed method pose a risk to consumers’ personal information? If so, is that risk outweighed by the benefit to consumers and businesses of using this method?

Section C. Invitation To Comment

You can file a comment online or on paper. For the Commission to consider your comment, we must receive it on or before September 30, 2014. Write “AgeCheq Application for Parental Consent Method, Project No. P–145410” on your comment. Your comment—including your name and your state—will be placed on the public record of this proceeding, including, to the extent practicable, on the Commission Web site, at http://www.ftc.gov/os/publiccomments.shtm. As a matter of discretion, the Commission tries to remove individuals’ home contact information from comments before placing them on the Commission Web site.

Because your comment will be made public, you are solely responsible for making sure that your comment doesn’t include any sensitive personal information, such as Social Security number, date of birth, driver’s license number or other state identification number or foreign country equivalent, passport number, financial account number, or credit or debit card number. You are also solely responsible for making sure that your comment does not include any sensitive health information, including medical records or other individually identifiable health information. In addition, do not include any “[t]rade secret or any commercial or financial information which is . . . privileged or confidential,” as provided in Section 6(f) of the FTC Act, 15 U.S.C. 46(f), and FTC Rule 4.10(a)(2). 16 CFR 4.10(a)(2). In particular, do not include competitively sensitive information such as costs, sales statistics, inventories, formulas, patterns, devices, manufacturing processes, or customer names.

If you want the Commission to give your comment confidential treatment, you must file it in paper form, with a request for confidential treatment, and follow the procedure explained in FTC Rule 4.9(c). 16 CFR 4.9(c). Your comment will be kept confidential only if the FTC General Counsel, in his or her sole discretion, grants your request in accordance with the law and the public interest.

Postal mail addressed to the Commission is subject to delay due to heightened security screening. As a result, we encourage you to submit your comments online. To make sure that the Commission considers your online comment, you must file it at https://ftcpublic.commentworks.com/ftc/coppagecheqapp, by following the instructions on the web-based form. If this Notice appears at http://www.regulations.gov/#/home, you also may file a comment through that Web site.

If you file your comment on paper, write “AgeCheq Application for Parental Consent Method, Project No. P–145410” on your comment and on the envelope, and mail it to the following address: Federal Trade Commission, Office of the Secretary, 600 Pennsylvania Avenue NW., Suite CC–5610 (Annex K), Washington, DC 20580, or deliver your comment to the following address: Federal Trade Commission, Office of the Secretary, Constitution Center, 400 7th Street SW., 5th Floor, Suite 5610 (Annex K), Washington, DC 20024. If possible, submit your paper comment to the Commission by courier or overnight service.

Visit the Commission Web site at http://www.ftc.gov to read this Notice and the news release describing it. The FTC Act and other laws that the Commission administers permit the collection of public comments to consider and use in this proceeding as appropriate. The Commission will consider all timely and responsive public comments that it receives on or before September 30, 2014. You can find more information, including routine uses permitted by the Privacy Act, in the Commission’s privacy policy, at http://www.ftc.gov/ftc/privacy.htm.

5 In particular, the written request for confidential treatment that accompanies the comment must include the factual and legal basis for the request, and must identify the specific portions of the comment to be withheld from the public record. See FTC Rule 4.9(c), 16 CFR 4.9(c). By direction of the Commission.

Janice Podoll Frankle, Acting Secretary.

[FR Doc. 2014–20645 Filed 8–28–14; 8:45 am]
BILLING CODE 6750–01–P

ENVIRONMENTAL PROTECTION AGENCY

40 CFR Part 52


Approval and Promulgation of Air Quality Implementation Plans;
Pennsylvania; Allegheny County’s Adoption of Control Techniques
Guidelines for Four Industry Categories for Control of Volatile
Organic Compound Emissions

AGENCY: Environmental Protection Agency (EPA).

ACTION: Proposed rule.

SUMMARY: The Environmental Protection Agency (EPA) is proposing to conditionally approve revisions to the Pennsylvania State Implementation Plan (SIP) submitted by the Commonwealth of Pennsylvania on behalf of the Allegheny County Health Department (ACHD). This SIP revision includes amendments to the ACHD Rules and Regulations, Article XXI, Air Pollution Control, and meets the requirement to adopt Reasonably Available Control Technology (RACT) for sources covered by EPA’s Control Techniques Guidelines (CTG) standards for the following categories: miscellaneous metal and/or plastic parts surface coating processes, automobile and light-duty truck assembly coatings, miscellaneous industrial adhesives, and fiberglass boat manufacturing materials. Upon review of the submittal, EPA found that the average monomer volatile organic compound (VOC) content limits were referenced but not included in the regulation for fiberglass boat manufacturing materials. ACHD has committed to revising the regulation and submitting the table of VOC content limits for fiberglass boat manufacturing materials to EPA in order to address specific RACT requirements for Allegheny County. EPA is, therefore, proposing conditional approval of the revisions to the Pennsylvania SIP in accordance with the requirements of the Clean Air Act (CAA).

DATES: Written comments must be received on or before September 29, 2014.

ADDRESSES: Submit your comments, identified by Docket ID Number EPA–