In FR Doc. 2016–24873 appearing on page 71369 in the Federal Register of Monday, October 17, the following corrections are made:

**Corrections**

1. On page 71369, in the third column, redesignate amendatory instructions 6 through 9 a through 10 and add new amendatory instruction 6 to read as follows:

   ■ 6. Amend §404.1592f by revising paragraph (a) to read as follows:

   §404.1592f How do we determine reinstated benefits?

   (a) If you meet the requirements for reinstatement under §404.1592c(a), we will then consider in which month to reinstate your entitlement. We will reinstate your entitlement with the earliest month, in the 12-month period that ends with the month before which you filed your request for reinstatement, that you would have met all of the requirements under §404.1592c(a) if you had filed your request for reinstatement in that month. Otherwise, you will be entitled to reinstated benefits beginning with the month in which you filed your request for such benefits if you did not perform substantial gainful activity in that month. If you performed substantial gainful activity in the month of filing, but are no longer able to perform substantial gainful activity, we will reinstate your benefits with the month after the month you filed your request for reinstatement. We cannot reinstate your entitlement for any month prior to January 2001.

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   Carolyn W. Colvin,
   Acting Commissioner of Social Security.

   [FR Doc. 2017–00076 Filed 1–19–17; 8:45 am]

   BILLING CODE 4191–02–P

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

**21 CFR Parts 73 and 74**

[Docket No. FDA–2016–F–0821]

**Listing of Color Additives Exempt From Certification; Titanium Dioxide and Listing of Color Additives Subject to Certification; [Phthalocyaninato (2-)] Copper; Confirmation of Effective Date**

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Final rule; confirmation of effective date.

**SUMMARY:** The Food and Drug Administration (FDA or we) is confirming the effective date of December 2, 2016, for the final rule that appeared in the Federal Register of November 1, 2016, and that amended the color additive regulations in §73.3126 (21 CFR 73.3126) and §74.3045 (21 CFR 74.3045) to provide for the safe use of titanium dioxide and [phthalocyaninato (2-)] copper to color orientation marks for intraocular lenses (IOLs). We are taking this action to ensure clarity that the effective date in the final rule remains December 2, 2016.

**DATES:** Effective date of final rule published in the Federal Register of November 1, 2016 (81 FR 75689), confirmed: December 2, 2016.

**FOR FURTHER INFORMATION CONTACT:** Laura A. Dye, Center for Food Safety and Applied Nutrition (HFS–265), Food and Drug Administration, 5001 Campus Dr., College Park, MD 20740–3835, 240–420–1275.

**SUPPLEMENTARY INFORMATION:** In the Federal Register of November 1, 2016 (81 FR 75689), we amended the color additive regulations in §73.3126 (21 CFR 73.3126) and §74.3045 (21 CFR 74.3045) to provide for the safe use of titanium dioxide and [phthalocyaninato (2-)] copper to color orientation marks for IOLs. The preamble to the final rule stated that persons who would be adversely affected by one or more provisions in the final rule could file electronic or written objections (81 FR 75689 at 75691). We also stated that the effective date of the final rule would be on December 2, 2016, unless a person properly files an objection or request for a hearing to review any provisions in the final rule (81 FR 75689). We explained that, to file an objection, a person must, among other things, specify with particularity the provision(s) of the regulation to which they object and the grounds for the objection (81 FR 75689 at 75691). Within each objection, a person also must specifically state whether he/she requests a hearing. We received no objections or requests for a hearing on the final rule that met these requirements. We received five general comments, including one that disagreed with the rule, but the comments did not meet the requirements to be considered an objection under 21 CFR 12.22(a)(3). Therefore, we find that the effective date of the final rule that published in the Federal Register of November 1, 2016, should be confirmed.

**List of Subjects**

21 CFR Part 73

Color additives, Cosmetics, Drugs, Medical devices.