quality of legal representation for children and youth in child welfare cases so the States and Tribes achieve the best safety, permanency and well-being outcomes for children and youth. This systems improvement model supports three research and demonstration sites, each involving a rigorous evaluation. Given the complexity of the models being implemented, considerable training, technical assistance, monitoring and support are necessary for each site to design and implement evaluation plans. Program expansion supplement funds will allow for an increased level of effort in conducting the evaluations in order to meet the requirements of the cooperative agreement. Additional training, technical assistance, and support to each research and demonstration site, coupled with more intensive monitoring of site specific evaluation efforts, will enhance the depth and rigor of all evaluation results.

The supplemental funding will also afford QIC-ChildRep the opportunity to provide new or modified technical assistance to assist States and Tribes in implementing the Administration on Children, Youth and Families' well-being framework in the context of the new requirements of the Child and Family Services Improvement and Innovation Act (Pub. L. 112–34).

Statutory Authority: Section 203 (42 U.S.C. 5113) of the Child Abuse Prevention and Treatment and Adoption Reform Act (CAPTA) of 1978, (Pub. L. 95–266), as amended.

Bryan Samuels,

Commissioner, Administration on Children, Youth and Families.

[FR Doc. 2012–26305 Filed 10–24–12; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[CFDA Number 93.674]

Announcement of the Award of a Single-Source Program Expansion Supplement Grant to the University of Oklahoma in Tulsa, OK

AGENCY: Children's Bureau, Administration on Children, Youth and Families, Administration for Children and Families, Health and Human Services.

ACTION: Announcement of the award of a single-source program expansion supplement grant to the University of Oklahoma, National Resource Center for Youth Development, in Tulsa, OK, to provide technical assistance to States to devise effective procedures and strategies to implement National Youth in Transition Database regulations effectively.

SUMMARY: The Administration for Children and Families (ACF), Children's Bureau (CB) announces the award of a single-source program expansion supplement in the amount of $103,685 to the University of Oklahoma, National Resource Center for Youth Development, Tulsa, OK, to support expanded technical assistance to address emerging technical assistance needs for States and Tribes as they seek to implement legislation and changing programs dedicated to former foster youth. The grantee is the recipient of a cooperative agreement to administer the National Resource Center for Youth Development (NRCYD). The grantee has been providing technical assistance services through a cooperative agreement since September 30, 2009, pursuant to the legislative authority of the Promoting Safe and Stable Families Program, Section 436(d), Title IV–B, subpart 2, of the Social Security Act (42 U.S.C. 629e).

DATES: September 30, 2012 through September 29, 2013.

FOR FURTHER INFORMATION CONTACT: Jan Shafer, Children's Bureau, 1250 Maryland Avenue SW., Washington, DC 20024. Telephone: 202–205–8172; Email: *jan.shafer@acf.hhs.gov*.

SUPPLEMENTARY INFORMATION: In February 2008, the National Youth in Transition Database (NYTD) final regulation was promulgated. NYTD requires States to begin collecting information from youth in foster care and young adults formerly in foster care every six months, beginning October 1, 2010. State representatives continue to identify implementation of NYTD as a significant challenge, particularly since it will require State agencies to remain in contact with youth who may no longer be receiving services from the agency. The implementation of NYTD will require the NRCYD to continue to provide additional technical assistance to States to implement this regulation effectively.

Additionally, many States see the implementation of NYTD as a method to engage youth and to strengthen youth involvement in services at the State and local level. This type of youth engagement work involves long-term systemic technical assistance. The single-source expansion supplement will allow the NRCYD to support these State initiatives over the long term.

Another significant development affecting the provision of services to youth and young adults was the passage of the Fostering Connections to Success and Increasing Adoptions Act of 2008, Public Law 110–351, signed into law October 7, 2008. Among other provisions, the law requires States to develop a transition plan for all youth emancipating from foster care and provides States and Tribes an option to receive Federal reimbursement under title IV–E of the Social Security Act to extend foster care to older youth until age 21. In addition, the law for the first time provided an opportunity for certain Tribes to receive direct funding for independent living services and education and training vouchers under the Chafee Foster Care Independence Program. The single-source program expansion supplement grant will allow the NRCYD to provide more intensive technical assistance and on-site consultation to States and Tribes to continue to assist them in implementing these provisions.

The supplemental funding will afford the National Resource Center for Youth Development the opportunity to provide new or modified technical assistance to assist States and Tribes in implementing the Administration on Children, Youth and Families' well-being framework in the context of the new requirements of the Child and Family Services Improvement and Innovation Act (Pub. L. 112–34).

Statutory Authority: Promoting Safe and Stable Families Program, § 436(d), Title IV–B, subpart 2, of the Social Security Act (42 U.S.C. 629e).

Bryan Samuels,

Commissioner, Administration on Children, Youth and Families.

[FR Doc. 2012–26304 Filed 10–24–12; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[CFDA Number 93.556]

Announcement of the Award of a Single-Source Program Expansion Supplement Grant to the Research Foundation of CUNY on Behalf of Hunter College School of Social Work, New York, NY

AGENCY: Children's Bureau, Administration on Children, Youth and Families, Administration for Children and Families, Health and Human Services.

ACTION: Announcement of the award of a single-source program expansion supplement grant to the Research Foundation of CUNY on behalf of Hunter College School of Social Work in New York, NY, to provide targeted technical assistance to Family Connections grantees.

SUMMARY: The Administration for Children and Families (ACF), Children's Bureau (CB) announces the award of a single-source program expansion supplement in the amount of $420,000 to the Research Foundation of CUNY on behalf of Hunter College School of Social Work, New York, NY, to provide targeted technical assistance to address continuing challenges in the field as child welfare programs work to implement the requirements of new legislation. The Research Foundation of CUNY on behalf of Hunter College is the recipient of a cooperative agreement to act as the administrator for the National Resource Center for Permanency and Family Connections (NRCPFC).

DATES: September 30, 2012 through September 29, 2013.

SUPPLEMENTARY INFORMATION: The supplemental funding will afford the National Resource Center on Permanency and Family Connections the opportunity to provide new or modified technical assistance to assist States and Tribes in implementing the Administration on Children, Youth and Families' well-being framework in the context of the new requirements of the Child and Family Services Improvement and Innovation Act (Pub. L. 112–34). In addition, the Fostering Connections to Success and Increasing Adoptions Act of 2008 (Pub. L. 110–351) provides for a discretionary matching grant program to implement projects in the areas of Kinship Navigator, Family Finding, Family Group Decision Making and Residential Family Treatment. The law also added a requirement at section 471(a)(29) that directs State foster care and adoption agencies (title IV–E agencies) to exercise due diligence to identify and notify all adult relatives of a child, within 30 days of the child's removal, of the relative's options to become a placement resource for the child. In total, the supplemental funding will allow the NRCPFC to do the following:

1. Provide focused technical assistance to Family Connections grantees.

2. Engage States that did not receive discretionary grants in on-site consultation regarding effectively involving relatives in child welfare practice.

3. Proactively transfer the knowledge developed under the discretionary grant program to States to assist in meeting new plan requirements.

The NRCPFC will increase technical assistance efforts to enhance the achievement of permanency by assisting agencies to better locate, notify and involve families and relatives in the engagement and planning process while maintaining awareness of confidentiality issues.

FOR FURTHER INFORMATION CONTACT: Jane Morgan, Children's Bureau, 1250 Maryland Avenue SW., Washington, DC 20024. Telephone: 202–205–8807; Email: *jane.morgan@acf.hhs.gov*.

Statutory Authority: Fostering Connections to Success and Increasing Adoptions Act of 2008 (Pub. L. 110–351).

Bryan Samuels,

Commissioner, Administration on Children, Youth and Families.

[FR Doc. 2012–26303 Filed 10–24–12; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2012–N–0007]

Generic Drug User Fee—Abbreviated New Drug Application, Prior Approval Supplement, and Drug Master File Fee Rates for Fiscal Year 2013

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the rate for the Abbreviated New Drug Application (ANDA), Prior Approval Supplement (PAS), and Drug Master File (DMF) fees related to the Generic Drug User Fee Program for fiscal year (FY) 2013. The Federal Food, Drug, and Cosmetic Act (the FD&C Act), as amended by the Generic Drug User Fee Amendments of 2012 (GDUFA), as further amended by the FDA User Fee Correction Act of 2012, authorizes FDA to assess and collect user fees for certain applications and supplements for human generic drug products, on applications in the backlog as of October 1, 2012, on finished dosage form (FDF) and active pharmaceutical ingredient (API) facilities, and on type II active pharmaceutical ingredient DMFs to be made available for reference. GDUFA directs FDA to establish each year the Generic Drug User Fee rates for the upcoming year. In the first year of GDUFA (FY 2013), some rates will be published in separate **Federal Register** notices because of the timing specified in the statute. Each year thereafter the GDUFA fee rates will be published 60 days before the start of the FY. This document establishes FY 2013 rates for an ANDA ($51,520), PAS ($25,760), and DMF ($21,340). These fees are effective on October 1, 2012, and will remain in effect through September 30, 2013.

FOR FURTHER INFORMATION CONTACT: David Miller, Office of Financial Management (HFA–100), Food and Drug Administration, 1350 Piccard Dr., PI50, rm. 210J, Rockville, MD 20850, 301–796–7103.

SUPPLEMENTARY INFORMATION:

I. Background

Sections 744A and 744B of the FD&C Act (21 U.S.C. 379j–41 and 379j–42), as added by GDUFA (Title III of the Food and Drug Administration Safety and Innovation Act (Pub. L. 112–144), which was signed by the President on July 9, 2012), as further amended by the FDA User Fee Correction Act of 2012 (Pub. L. 112–193) (signed by the President on October 5, 2012), establish fees associated with human generic drug products. Fees are assessed on the following: (1) Certain applications in the backlog as of October 1, 2012; (2) certain types of applications and supplements for human generic drug products; (3) certain facilities where APIs and FDFs are produced; and (4) certain DMFs associated with human generic drug products (section 744B(a) of the FD&C Act). This notice will focus on the ANDA, PAS, and DMF fees.

II. Fee Revenue Amount for FY 2013

The total fee revenue amount for FY 2013 is $299,000,000, as set in the statute. GDUFA directs FDA to use the yearly revenue amount as a starting point to set the fee rates for each fee type. GDUFA states that the backlog fee will make up $50,000,000 of the total revenue collected for FY 2013. Therefore, the rest of the fees will make up a percentage of the remaining $249,000,000 of the total revenue. For more information about GDUFA, please refer to the FDA Web site (*http://www.fda.gov/gdufa*). The ANDA, PAS, and DMF fee calculations for FY 2013 are described in this document.

III. ANDA and PAS Fees

Under GDUFA, the ANDA and PAS fees are owed by each applicant that submits, on or after October 1, 2012, an ANDA or a PAS. These fees are due on the date of submission of the ANDA or PAS or 30 days after the publication date of this notice, whichever is later. Section 744B(b)(2)(B) specifies that the