PLEASE READ INSTRUCTIONS CAREFULLY. Be sure to indicate any changes in your legal name or actual location in item 4, and any changes in your mailing address in item 6. Print all entries and make all corrections in red ink, if possible. Enter your phone number in item 8.3 and the phone number of your actual location in item 4.1. Sign the form and return to FDA. After validation, you will receive your Official Registration for the 6. MAILING ADDRESS OF REPORTING OFFICIAL (Include institution name if 5. OTHER NAMES USED AT THIS LOCATION (Include trade name, doing-business-as, previous names, and other firms co-located. If applicable, include registration 4.1 PHONE 865-524-3074 state, country, and post office code) 4. LEGAL NAME AND LOCATION (Include legal name, number and street, city, 8.1 TYPED NAME Martha S. Cox, Chief Quality Officer 8. REPORTING OFFICIAL'S SIGNATURE 7.2 PHONE 7.1 E-MAIL ADDRESS state, and zip code) 7. U.S. AGENT (Include name, institution name if applicable, number and street, city, applicable, number and street, city, state, country, and post office code) ENTER ALL CHANGES IN RED INK AND CIRCLE. 8.3 PHONE 865-524-3074 x668 8.2 E-MAIL ADDRESS mcox@medicblood.org ensuing year. MEDIC, Inc. Medic Regional Blood Center Medic, Inc. ATTN: Martha S. Cox, Chief Quality Officer Medic, Inc. Knoxville, TN 37921-6702 Knoxville, TN 37921-6702 1601 Ailor Avenue 1601 Ailor Avenue **BLOOD ESTABLISHMENT REGISTRATION AND PRODUCT LISTING** DEPARTMENT OF HEALTH AND HUMAN SERVICES PUBLIC HEALTH SERVICE FOOD AND DRUG ADMINISTRATION Curtus 8.4 DATE 1/19/2017 result in a fine of up to \$1,000 or imprisonment up to one year or both, pursuant to Section 303(a) of the Act (Title 21, United States Code 33.3(a)). This form is authorized by Sections 510(b), (j) and 704 of the Federal Food, Drug, and Cosmetic Act (Title 21, United States Code 360(b), (j) and 374). Failure to report this information is a violation of Section 301(f) and (p) of the Act (Title 21, United States Code 331(f) and (p)) and can OTHER Cryoprecipitated AHF Pooled BLOOD BANK REAGENTS BLOOD PRODUCTS FOR DIAGNOSTIC USE RECOVERED PLASMA SOURCE PLASMA SOURCE LEUKOCYTES FRESH FROZEN PLASMA PLASMA CRYOPRECIPITATE REDUCED PLASMA PLATELETS RBC REJUVENATED WHOLE BLOOD THERAPEUTIC EXCHANGE PLASMA LIQUID PLASMA CRYOPRECIPITATED AHF RBC REJUVENATED FROZEN RBC FROZEN RED BLOOD CELLS (RBC) LEUKOCYTES/GRANULOCYTES RBC REJUVENATED DEGLYCEROLIZED RBC DEGLYCEROLIZED 11. PRODUCTS ALLOGENEIC .9 OTHER (Specify) : .7 STATE .6 U.S. MILITARY .5 FEDERAL (non-military) .4 COOPERATIVE ASSOCIATION .3 CORPORATION .2 PARTNERSHIP .1 SINGLE PROPRIETORSHIP TYPE OF OWNERSHIP .8 \bigsqcup COUNTY/MUNICIPAL/HOSPITAL AUTHORITY Platelet Pheresis (automated) AUTOLOGOUS × 2. U.S. LICENSE NUMBER 1. REGISTRATION NUMBER profit_ FEI: CFN: 1077605 DIRECTED 1077605 non-profit 19 12 \vec{b} 20 18 77 6 햐 ಘ コ 6 COLLECT Ξ × × MANUAL APHERESIS 10. TYPE ESTABLISHMENT (Check all boxes that describe routine or autologous operations.) <u>8</u> .3 CHANGE IN INFORMATION .2 INITIAL REGISTRATION .1 ANNUAL REGISTRATION 3. REASON FOR SUBMISSION 10 OTHER (Specify) : .6 COMPONENT PREPARATION FACILITY .7 COLLECTION FACILITY .8 DISTRIBUTION CENTER .5 HOSPITAL TRANSFUSION SERVICE .1 🔽 COMMUNITY (NON-HOSPITAL) BLOOD BANK .9 BROKER/WAREHOUSE .3 PLASMAPHERESIS CENTER .4 PRODUCT TESTING LABORATORY .2 HOSPITAL BLOOD BANK AUTOMATED APHERESIS × × × (ω -NOT APPROVED FOR MEDICARE REIMBURSEMENT INDEPENDENT APPROVED FOR MEDICARE REIMBURSEMENT ASSOCIATED W/ COMMUNITY or HOSPITAL BLOOD BANK PREPARE <u>;</u> × Ħ × × × × × × × REDUCED (5) × × × × VALIDATED BY FDA: 23-DEC-2016 PRINTED BY FDA: DISTRICT OFFICE: IRRADIATED × × × × × 6) × × × U.S. LICENSE NUMBER OF PARENT FIRM 688 FOR FDA USE ONLY DONOR RETESTED $\widehat{\Xi}$ 18-JAN-2017 New Orleans TEST (B) STORE and DISTRIBUTE to OTHERS (0. × × × × × × × × × × × ×