

Health and Human Services Agency California Department of Public Health



Erica Pan, MD, MPH
Director and State Public Health Officer

Gavin Newsom
Governor

DATE: July 30, 2025

TO: Local Health Officers

County Health Executives Association of California (CHEAC) Members

Immunization Coordinators

Local Health Department Fiscal Staff

Receiving Immunization Program Local Assistance Grants

FROM: Maria E. Volk, MPA, Assistant Branch Chief

Immunization Branch

marie a. Volh

SUBJECT:

- 1.) Immunization Local Assistance Funding, Fiscal Year 2025-26
- 2.) Federal Compliance Requirements of the Immunization Grant Fiscal Year 2025-26 Acknowledgement and Subrecipient Information Form
- 3.) Fiscal Year 2025-26 Immunization Agreement Invoice Submission Deadlines and Budget Modification Guidelines

GRANT AGREEMENT FUNDING ANNOUNCEMENT/RELEASE

The California Department of Public Health (CDPH), Immunization Branch, is pleased to release the Federal Grant subaward notification to Local Health Department (LHD) grantees for FY 2025-26. CDPH has authority to grant funds for the Project under Health and Safety Code, Section 120325-120380, which requires immunizations against childhood diseases prior to school admittance. The purpose of this grant is to assist LHDs in preventing and controlling vaccine-preventable diseases in the local health jurisdiction (LHJ).

RELATED STATUTES

California Health & Safety Code sections:

 120130 requires the Local Health Officer to properly report to CDPH those diseases listed as reportable, which include vaccine-preventable diseases.



Local Health Officers Immunization Coordinators Receiving Immunization Program Local Assistance Grants Page 2 July 30, 2025

- 120175 requires the Local Health Officer to take measures as may be necessary to prevent the spread or occurrence of additional cases of reportable diseases (which includes reportable vaccine-preventable diseases).
- 120350 requires Local Health Officers to organize and maintain a program to make available the immunizations required for admittance to childcare facilities and schools.

SERVICES TO BE PERFORMED BY THE GRANTEE

The Grantee is to implement activities to:

- Assess and improve coverage levels in the jurisdiction of all vaccines recommended by the Advisory Committee on Immunization Practices (ACIP) to protect the population.
- Detect, report, and control vaccine-preventable diseases in the jurisdiction.
- For detailed scope of work objectives and activities, please refer to the enclosed revised Scope of Work. The CDPH and award subrecipients will make agreed-upon changes to the Scope of Work on an as-needed basis. These changes will be made in writing but will not require a formal amended agreement.

OVERVIEW, GRANT TERMS, AND FUNDING

The immunization award for fiscal year 2025-26 is reduced from last year. Please read the award information below on the Federal Grant Award. The effective date for this award is July 1, 2025. Similar to prior years, the availability of federal local assistance funding is dependent upon funds received from CDC and, at CDPH's discretion.

IMMUNIZATION FEDERAL AWARD

- Federal Grant Award No.: 1 NH23IP922690-01-00
- Award Issue Date: 07/03/2025
- Catalog of Federal Domestic Assistance (CFDA) Title: Immunization Cooperative Agreements
- Catalog of Federal Domestic Assistance (CFDA) No.: 93.268
- Data Universal Numbering Systems (DUNS) No.: 7991506150000
- Unique Entity Identifier (UEI) No.: KD2JSY6LNMW7
- Total Federal Award to Date: \$32,190,571
- Amount Made Available for Local Assistance Subrecipient Awards: \$5,917,955
- Year 4 Budget, FY 2025-26: 100% Prevention and Public Health Funds (PPHF)

STATE GENERAL FUND

State General Funds are included in Local Assistance funding for support of your continuing Local Health Department efforts to prevent influenza.

Local Health Officers Immunization Coordinators Receiving Immunization Program Local Assistance Grants Page 3 July 30, 2025

ELIGIBLE FOR LOCAL ASSISTANCE:

The Immunization Branch has determined that the following 60 LHDs and two non-profit organizations are eligible to apply for available funding for their Local Immunization Program, which supports the State's objectives to control vaccine-preventable diseases.

County of Alameda	County of Madera	County of San Joaquin
County of Alpine	County of Marin	County of San Luis Obispo
County of Amador	County of Mariposa	County of San Mateo
City of Berkeley	County of Mendocino	County of Santa Barbara
County of Butte	County of Merced	County of Santa Clara
County of Calaveras	County of Modoc	County of Santa Cruz
County of Colusa	County of Mono	County of Shasta
County of Contra Costa	County of Monterey	County of Sierra
County of Del Norte	County of Napa	County of Siskiyou
County of El Dorado	County of Nevada	County of Solano
County of Fresno	County of Orange	County of Sonoma
County of Glenn	City of Pasadena	County of Stanislaus
County of Humboldt	County of Placer	County of Sutter
County of Imperial	County of Plumas	County of Tehama
County of Inyo	County of Riverside	County of Trinity
County of Kern	County of Sacramento	County of Tulare
County of Kings	County of San Benito	County of Tuolumne
County of Lake	County of San Bernardino	County of Ventura
County of Lassen	County of San Diego	County of Yolo
City of Long Beach	City & County of San Francisco	County of Yuba

CA Immunization Coalition CA Primary Care Association

Please complete, sign, and return the following to <u>izb.admin@cdph.ca.gov</u> as soon as possible or no later than August 13, 2025.

- Federal Compliance Requirements of the Immunization Grant No. 1 NH23IP922690-01-00.
- Subrecipient Information Form is being requested to ensure we have the most current information on file.
- Exhibit B Budget worksheet, please complete and submit the award worksheets for the Vaccine Preventable Disease (VPD) Prevention grant AND the worksheet for the State General Fund (SGF) grant.

Local Health Officers Immunization Coordinators Receiving Immunization Program Local Assistance Grants Page 4 July 30, 2025

FY 2025-26 Immunization Agreement Invoice Submission Deadlines and Budget Modification Guidelines

Invoices are due on a quarterly basis. The final invoice for the fiscal year is due no more than sixty (60) calendar days from June 30.

Quarter	Invoice Service Period	Invoice Due date
Quarter 1	July 1, 2025 – September 30, 2025	11/14/2025
Quarter 2	October 1, 2025 – December 31, 2025	2/13/2026
Quarter 3	January 1, 2026 – March 31, 2026	5/15/2026
Quarter 4	April 1, 2026 – June 30, 2026	8/28/2026

Budget modification requests are required when shifting/moving funds from one budget line-item to another. Budget line-item shifts do not require a formal grant agreement amendment and can be agreed upon between CDPH and the subrecipient. Budget Modification Requests must be submitted and approved prior to submitting the corresponding invoice. (See enclosed Budget Modification Request).

Thank you.

Enclosures: Award-6 NH23IP922612-05-05 GTC Non-Research

IZB Grant 2025-26 SOW

IZB Grant 2025-26 Required Activities SOW

Federal Compliance & Subrecipient Information Form FY2025-26

FY2025-26_Budget by Grant

Budget Modification Request Form FY2025-26

CC:

State Immunization Branch Field Representatives

Noemi Marin, CDPH, Immunization Branch

Raquel Espinoza, CDPH, Immunization Branch

Roland Rafol, CDPH, Immunization Branch

Rodolfo Ruiz, CDPH, Immunization Branch

Kae Saechao, CDPH, Immunization Branch

From: Espinoza, Raquel@CDPH < Raquel. Espinoza@cdph.ca.gov>

Sent: Wednesday, August 6, 2025 9:56 AM

To: VanEarden, Melenie < Melenie. VanEarden@dph.sbcounty.gov>; Guerrero, Kathy

<Kathy.Guerrero@dph.sbcounty.gov>; Ibrahim, Diana <Diana.Ibrahim@dph.sbcounty.gov>; Hernandez, Jenny

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<dramirez@dph.sbcounty.gov>; Chandler, Trent <Trent.Chandler@dph.sbcounty.gov>

Cc: Dugas, Joshua <Joshua.Dugas@dph.sbcounty.gov>; Wang, Sharon <Sharon.Wang@dph.sbcounty.gov>

Subject: Immunization funding allocation information FY 25/26

Importance: High

CAUTION: This email originated from outside of the organization. Do not click links or open attachments unless you can confirm the sender and know the content is safe.

Immunization Coordinator,

Below is this year's immunization funding allocation information.

County: San Bernardino

CDPH Grant #: 22-11081 FY2025-26 Allocation: 427,221.00

Grant Award Award Amount
Federal VPD Grant 283,504.00
State General Fund (SGF) Grant 143,717.00

Please complete, sign, and return the following no later than August 13, 2025.

- Federal Compliance & Subrecipient Information Form
- Budget worksheets
 - Vaccine Preventable Disease (VPD) Prevention grant
 - State General Fund (SGF) grant

As a reminder, budget modification requests are required when shifting/moving funds from one budget line-item to another. Budget line-item shifts do not require a formal grant agreement amendment. Budget Modification Requests must be submitted and approved prior to submitting the corresponding invoice. Modification form attached.

Invoice Service Period	Invoice Due date
July 1, 2025 – September 30, 2025	11/14/2025
October 1, 2025 – December 31, 2025	2/13/2026
January 1, 2026 – March 31, 2026	5/15/2026
April 1, 2026 – June 30, 2026	8/28/2026
	July 1, 2025 – September 30, 2025 October 1, 2025 – December 31, 2025 January 1, 2026 – March 31, 2026

Let me know if you have any questions or concerns.

Thank you,

Raquel Espinoza

Immunization Branch
California Department of Public Health (CDPH)
850 Marina Bay Parkway, Building P, 2nd Floor
Richmond, CA 94804

Raquel.espinoza@cdph.ca.gov

W: (279) 217-1198













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CDPH Immunization Branch Local Assistance Federal Fund VPD Grant

County: San Bernardino CDPH Grant #: 22-11081

Year 4 - Budget Detail Fiscal Year: 07/01/2025 - 06/30/2026

	% of time or hours on project	Monthly salary range or hourly		
I. Personnel		rate		Total
Communicable Disease Investiagor II Health Services Assistant Office Specialist PH Program Manager Public Health Nurse Senior Accountant / Auditor Senior Office Assistant Supervising Health Svcs Assistant Supervising Office Assistant	40% 40% 15% 5% 45% 5% 5% 5% 25%	\$ 4,881.85 \$ 5,329.80 \$ 11,773.41 \$ 13,102.23 \$ 8,338.60 \$ 4,967.59 \$ 5,192.66	\$ \$ \$ \$	32,192.17 23,432.87 9,593.63 7,064.04 70,752.06 5,003.16 2,980.56 3,115.60 16,682.87
			\$ \$	-
Total Personnel Expenses			\$	170,816.95
II. Fringe Benefits (40.92% of Personnel)			\$	69,898.30
III. Operating Expenses or General Expenses			\$	389.17
Communications			\$	389.17
Office/Clinic Supplies			\$	-
Postage			\$ \$ \$	-
Special Department Expense			\$	-
IV. Equipment Expenses			\$	-
			\$	-
V. Travel Expenses			\$	_
In-State Travel			\$	-
Out-of-State Travel (The Grantee shall be reimbursed for the actual claime	ed and invoiced)		\$	-
VI. Subgrantee Expenses (if any)(Name of Subgrantee)			\$	-
VII. Indirect Costs (Approved% of total Personnel C	osts or total Direct	Costs)	\$	42,399.58
☒ Approved 17.614% of total Personnel Costs☐ Approved% of total Direct Costs				
VIII. Total Expenses For Main Grant			\$	283,504.00

CDPH Immunization Branch Local Assistance State General Fund Grant

County: San Bernardino

CDPH Grant #: 22-11081

Year 4 - Budget Detail Fiscal Year: 07/01/2025 - 06/30/2026

I. Personnel	% of time or hours on project	Monthly salary range or hourly rate		Total
Communicable Disease Investiagor II Health Services Assistant Office Specialist PH Program Manager Public Health Nurse Senior Accountant / Auditor Senior Office Assistant Supervising Health Svcs Assistant Supervising Office Assistant	20% 20% 7% 3% 20% 5% 5% 3% 10%	\$ 6,706.70 \$ 4,881.85 \$ 5,329.80 \$ 11,773.41 \$ 13,102.23 \$ 8,338.60 \$ 4,967.59 \$ 5,192.66	\$ \$ \$ \$ \$ \$ \$ \$ \$ \$ \$	16,096.08 11,716.43 4,477.03 4,238.43 31,445.36 5,003.16 2,980.56 1,869.36 6,673.15
Total Personnel Expenses			\$	84,499.55
II. Fringe Benefits (40.92% of Personnel)			\$	34,577.22
III. Operating Expenses or General Expenses Communications Office/Clinic Supplies Postage Special Department Expense			\$ \$ \$ \$	3,666.05 2,116.05 500.00 50.00 1,000.00
IV. Equipment Expenses			\$	-
V. Travel Expenses In-State Travel Out-of-State Travel (The Grantee shall be reimbursed for the actual claimed	and invoiced)		\$ \$ \$	- - -
VI. Subgrantee Expenses (if any) (Name of Subgrantee)			\$	-
VII. Indirect Costs (Approved 17.614% of total Personne	el Costs or total Dir	rect Costs)	\$	20,974.18
 ☑ Approved 17.614% of total Personnel Costs ☐ Approved% of total Direct Costs 		,	·	
VIII. Total Expenses For Main Grant			\$	143,717.00

Purpose

The purpose of this grant is to assist local health departments (LHDs) in preventing and controlling vaccine-preventable diseases in the local health jurisdiction (LHJ).

Related Statutes

California Health & Safety Code sections:

- 120130 requires the Local Health Officer to properly report to CDPH those diseases listed as reportable, which include vaccine-preventable diseases.
- 120175 requires the Local Health Officer to take measures as may be necessary to prevent the spread or occurrence of additional cases of reportable diseases (which includes reportable vaccine-preventable diseases).
- 120350 requires Local Health Officers to organize and maintain a program to make available the immunizations required for admittance to childcare facilities and schools.

Services to be Performed by the Grantee

The Grantee is to implement activities to:

- Assess and improve coverage levels in the jurisdiction of all vaccines recommended by the Advisory Committee on Immunization Practices (ACIP) to protect the population.
- Detect, report, and control vaccine-preventable diseases in the jurisdiction.

The LHD must agree to the following inclusive objectives and conduct the following activities. Many of the services to be performed are also conditions for federal funding of the CDPH Immunization Branch (IZB) and/or statutory requirements of State and LHDs. The level of local assistance grant funding to be awarded is not represented as sufficient for support of all the required activities; a significant amount of local support and funding is expected. Local assistance grant funds must not be used to supplant (i.e., replace) local funds currently being expended for immunization services and activities.

Grantee agrees to assign the responsibility of monitoring each program component:

1) Vaccine Accountability and Management; 2) Access to and Utilization of Quality Immunization Services; 3) California Immunization Registry (CAIR); 4) Perinatal Hepatitis B Prevention; 5) Education, Information, Training, and Partnerships; 6) Prevention, Surveillance and Control of Vaccine Preventable Disease (VPD); 7) Childcare and School Immunization Entry Requirements; and 8) Influenza.

Grantee will monitor grant fund expenditures to maximize the utilization of the funding for achieving the goals and objectives. Grant invoices shall be reviewed and submitted quarterly to the CDPH Immunization Branch.

The Immunization Coordinator is required to participate in meetings, webinars, and conference calls as requested by the CDPH Immunization Branch including, but not limited to, the CDPH Immunization Branch's Immunization Coordinators' Meeting, New Immunization Coordinator Orientation (offered annually and required for all new Immunization Coordinators), regional coordinators' meetings, and conference calls related to influenza, outbreak control, perinatal hepatitis B, changes in policies and procedures, and other important issues.

Area 1. Vaccine Accountability and Management

Goal 1.1: Maintain viability of IZB-supplied vaccine to ensure vaccine effectiveness and reduce vaccine waste.				
Required Activities	Performance Measures			
Goal 1.1 Activity a: Annually, make sure all relevant staff within LHD-operated clinics (routine mass vaccination, or special immunization outreach) are properly trained on current policies and procedures for proper vaccine storage and handling outlined in each participation agreement/addendum for the receipt of IZB-supplied vaccines (317, Vaccines for Children [VFC], state general fund).	 Updated Vaccine Management Plans for each LHD facility. Completed EZIZ Lessons for Key Practice Staff. Completed training logs (training date, topics, methods, and list of attendees). 			
Goal 1.1 Activity b: Develop and implement a training plan for provider facilities outside LHDs receiving IZB supplied doses (state or 317 Outbreak). Focus the plan on proper vaccine management, vaccine storage and handling requirements, and administration prior to the distribution of IZB-supplied vaccines. Goal 1.1 Activity c: Develop and implement a plan to verify that providers administering 317 Outbreak and state general fund immunizations outside the LHDs adhere to policies for vaccine management. Conduct Quality Assurance verifications (such as random temperature log review, on-site vaccination clinic assessments, review of vaccine transport logs, and when indicated, review of vaccine losses, etc.) at least every other year, in a sample of sites receiving SGF vaccines, and for outbreak, within the outbreak effort, in a sample of outside partners.	 Training plan developed and implemented. Number of completed trainings. Completed training logs (training date, topics, methods, and list of attendees). Training packet completed and available. Number of signed Vaccine Management Plans received and reviewed. Developed and implemented Quality Assurance Plan. Completion of Mass Vaccination Hourly Temperature Logs/Electronic Data Files. Temperature Documentation on CDPH provided Logs for all IZB-supplied vaccines/Electronic Temperature Files. Percentage of sites receiving Quality Assurance verifications (minimum sample of 10% of sites receiving vaccines). Number of Completed Quality Assurance verifications. 			

Goal 1.2: Facilitate compliance with current protocols, policies, and procedures for vaccine accountability for LHD facilities and partners that receive IZB-supplied vaccine.				
Activity	Performance Measures			
Goal 1.2 Activity a: Make sure all relevant staff involved in vaccine ordering, management, and accountability activities within local health department-operated clinics adhere to all program requirements as outlined in the VFC/317 Provider Participation Agreements and Addendums. Complete annual VFC/317 program recertification.	Completed annual program recertification and corresponding educational lessons for all key practice staff.			
Goal 1.2 Activity b: Promote adherence to eligibility guidelines corresponding to VFC, Section 317, and state general fund vaccines. Upon release of the Immunization Branch's Vaccine Eligibility Guidelines, IMM-1142, disseminate guidance to all relevant staff involved in vaccine ordering, management, and accountability activities within local health department operated pediatric and adult immunization clinics.	Documentation of provided guidance.			
Goal 1.2 Activity c: Verify that processes are in place such that IZB-supplied (317, VFC, state) vaccines are administered to eligible individuals following outlined eligibility guidelines for each vaccine funding source.	Updated LHD protocols, inclusive of eligibility guidelines, for each vaccine funding source.			
Goal 1.2 Activity d: Comply with federal policies regarding vaccine distribution. Publicly funded VFC and 317 vaccines must be distributed directly to the location at which the provider will administer the vaccines.	Documentation of procedures.			

Area 2. Access to and Utilization of Quality Immunization Services

Goal 2.1: Improve access to and receipt of all ACIP-recommended immunizations, especially for low income and underserved community members.			
Required Activities Performance Measures			
Goal 2.1 Activity a : Provide LHD resource and referral lists to other programs that connect patients to services.	Referral list completed and updated on an annual basis.		
Goal 2.1 Activity b: For all <i>LHD facilities</i> that are VFC providers, participate in and support provider compliance and quality improvement visits in conjunction with the CDPH	Number of LHD clinics with corrective actions that were all completed within the specified time frame on the VFC Compliance Visit Report.		

Immunization Branch. Assist with the	2. Number of LHD IQIP visits.
implementation of corrective action plans,	
strategies to reduce missed opportunities for	
vaccination, and linkage/referral to medical	
homes.	

Area 3. California Immunization Registry (CAIR)¹

Goal 3.1 Promote and optimize ² the use of CAIR in the jurisdiction				
Required Activities	Performance Measures			
Goal 3.1 Activity a: Ensure all IZB-supplied vaccine doses administered by LHD or partners, including influenza doses, are entered into CAIR.	 Number of LHD clinics participating in CAIR/ number of LHD clinics. Percentage of LHD clinic doses entered into the registry within 14 days. Number of state flu doses entered by end of 			
Note: Any entity that is authorized to use the California Immunization Registry must adhere to the regulations outlined in California Health and Safety Code 120440 which governs immunization registry data use. For more information, please visit HSC 120440 (ca.gov).	flu season/number of state flu doses administered. 4. CAIR ID list submitted to CDPH.			
Goal 3.1 Activity b: For LHDs with primary care clinics, should use the Manage Patient Status functionality to remove inactive patients at least once a year. Training is available for staff who need assistance with the process.	Inactive patients marked as inactive in CAIR.			
Goal 3.1 Activity c: In LHD primary care clinics, utilize CAIR data to identify and improve low or lagging infant or adolescent vaccination coverage levels.	Low infant or adolescent CAIR coverage rate identified and improved.			

Goal 3.2: Connect local Immunization Information Systems (IIS) to CAIR (for San Joaquin County only)		
Required Activities	Performance Measures	
Goal 3.2 Activity a: Maintain ongoing	Ongoing data sharing continues.	
electronic data sharing with CAIR2.		

¹ CAIR refers to the statewide system that will connect CAIR2 with the San Diego Immunization Registry and Healthy

² If have EHR, move from manual data entry to data exchange (upload from EHR) to bidirectional data exchange. See https://www.cdph.ca.gov/Programs/CID/DCDC/CAIR/CDPH%20Document%20Library/IMM-1266.pdf and https://www.cdph.ca.gov/Programs/CID/DCDC/CAIR/CDPH%20Document%20Library/IMM-1260.pdf

Area 4. Perinatal Hepatitis B Prevention

Goal 4.1: Reduce the incidence of perinat	al hepatitis B virus (HBV) in	ifection in the
jurisdiction.		

Note: Coordinate perinatal HBV prevention efforts with your LHD's Maternal Child and Adolescent Health (MCAH) program, as activities 4.1a-4.1c may also help fulfill Title V requirements and MCAH Scope of Work Activities.

MCAH Scope of Work Activities.	
Required Activities	Performance Measures
Goal 4.1 Activity a: Educate medical providers and hospital staff about the screening, care, and reporting of pregnant persons who test positive for hepatitis B and their infants according to the guidance outlined below: Guidance for Prenatal Providers Guidance for Labor and Delivery Hospitals Guidance for Pediatric Providers	 Percentage of HBsAg-positive pregnant women identified in the reporting period with an HBV DNA test result during pregnancy. Percentage of infants born to HBsAg-positive mothers in the reporting period who received PEP according to ACIP recommendations.
Goal 4.1 Activity b: Educate identified HBsAg-positive pregnant persons about their HBV status and provide the appropriate information on prevention of perinatal hepatitis B transmission, based on current ACIP recommendations and the guidance outlined in the CDPH Perinatal Hepatitis B Prevention Program Coordinator Handbook Note: Even if you had no cases in the previous grant period you are still expected to review the Handbook internally so that you are prepared to reach out to patients in the future.	Percentage of HBsAg-positive pregnant women identified in the reporting period who were reported and contacted prior to delivery.
Goal 4.1 Activity c: Collect and submit requested data to CDPH on pregnant persons who test positive for hepatitis B and their infants according to the guidance outlined below: Perinatal Hepatitis B Prevention Program Coordinator Handbook	 Percentage of PEP errors in the reporting period with completed LHJ follow-up (defined as submission of a PEP Error Reporting Form to CDPH PHPP). Percentage of infants born to pregnant persons with hepatitis B who complete the HBV vaccine series by 12 months of age. Percentage of infants born to pregnant persons with hepatitis B who complete PVS testing by 24 months of age. Percentage of infants closed to case management with complete information within 24 months.

Area 5. Education, Information, Training, and Partnerships

Goal 5.1: Provide and/or promote educational activities a	and information to health care
providers, schools and childcare centers, and other imm	unization stakeholders to promote
best practices for immunizations and the importance of timely vaccinations.	

best practices for immunizations and the importance of timely vaccinations.	
Required Activities	Performance Measures
Goal 5.1 Activity a: Based on local priorities and resources, disseminate print and/or electronic communications among providers, school, general public and other immunization stakeholders in their jurisdiction.	 Summary of efforts conducted to distribute materials in print or electronically to immunization stakeholders. Target date for completion of summary.
Note: Depending on funding, CDPH offers select hard-copy materials that immunization coordinators can order from the Branch Store at izcoordinators.org. CDPH will inform LHDs on communication activities from the Immunization Branch (e.g., print materials available; electronic communications and resources for VFC providers, schools, pharmacies, and community-based organizations/other stakeholders; and traditional media/social media activities to reach the general public). LHDs may supplement any gaps in communication with local efforts. Contact the Information & Education Section if you would like to learn more about the Immunization Branch's centralized communication vehicles and activities.	

Goal 5.2: Develop partnerships and collaborative activities in order to expand immunization services, promote best practices and improve coverage rates among children, adolescent and adults.	
Required Activities	Performance Measures
All activities for this goal are Supplemental Activities.	

Area 6. Prevention, Surveillance and Control of Vaccine Preventable Disease (VPD)

Goal 6.1: Conduct surveillance to identify VPD cases and/or outbreaks, and implement recommended prevention and control activities.	
Required Activities	Performance Measures
Goal 6.1 Activity a: Ensure that appropriate	Percentage of measles PCR positive
clinical specimens are tested, and relevant	specimens submitted for molecular
	characterization.

epidemiologic information is collected for VPDs requiring immediate public health action.	2. Percentage of Neisseria meningitidis positive specimens/isolates submitted for molecular characterization.
Goal 6.1 Activity b: Implement appropriate public health activities for the control and prevention of cases and/or outbreaks of VPDs in accordance with CDPH recommendations including obtaining vaccine and taking efforts to vaccinate susceptible individuals, as appropriate.	 Quarterly review of Quicksheets with applicable staff completed. Percentage of infant pertussis cases <4 months of age with documentation of mother's prenatal care provider information (name and city of prenatal care provider). Percentage of infant pertussis cases <4 months of age for whom maternal Tdap status is known. Percentage of providers reached³ who provided prenatal care to a woman whose infant developed pertussis. Reaching the provider is only required if the woman never got Tdap during this pregnancy or did not receive Tdap at 27-36 weeks gestation.⁴ Completed outbreak response request⁵ with plan for doses and target population (as appropriate).

Goal 6.2: Collect and submit requested data to CDPH on VPD cases and outbreaks.	
Required Activities	Performance Measures
Goal 6.2 Activity a: Classify VPD cases with appropriate CSTE resolution status and submit case and outbreak data to CDPH per instructions listed here: ReportingGuidanceForLHJs	 Percentage of VPD cases with appropriate resolution status assigned, as per CSTE case definition. Average percentage correct of all VPDs reported. Percentage of measles cases reported immediately to CDPH. Percentage of confirmed hepatitis A cases for whom hepatitis A risk factors are known. Percent of meningococcal disease cases in high school and college students reported immediately to CDPH. Percentage of meningococcal disease cases
	aged 14-24 years for whom high school or college attendance status is known.

³ Sending a letter re: standard of care is the minimum acceptable communication, with copy to your LHD Maternal Child and Adolescent Health (MCAH) program. See *Template Letter for Prenatal Care Providers with Pregnant Patients that did not Receive Prenatal Tdap Appropriately and Infants Developed Pertussis*

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⁴ Please note: If a practice decides to modify or reduce the number of prenatal care visits due to the COVID-19 pandemic, ACOG encourages clinicians to include recommended maternal immunizations (influenza and Tdap) during remaining inperson appointments, even if that means immunizations will be administered outside of the typically recommended weeks of gestation. Please make note of any clinic modifications in the notes field in the maternal immunizations section in CalREDIE.

⁵ The Immunization Branch provides a form for requesting vaccine from CDPH.

Percentage of case reports submitted to
CDPH via an electronic communicable
disease reporting system (CalREDIE or
other) in the recommended timeframe.

Area 7. Childcare and School Immunization Entry Requirements

Goal 7.1: Decrease the proportion of pupils who are overdue for required immunizations or	
admitted conditionally. Required Activities	Performance Measures
Goal 7.1 Activity a: Provide guidance, training, and support for compliance with entry immunization requirements by all childcare centers and schools within the jurisdiction.	Percentage of schools with kindergarteners in the jurisdiction that have completed the annual immunization assessment.
Goal 7.1 Activity b: At least annually, visit schools (either in-person or virtually) with 10 or more kindergarteners that reported > 10% were either conditionally admitted or overdue for required immunization; provide guidance and support follow-up until these students are up to date.	 Number and percentage of schools that meet the definition of "targeted schools" ⁶ Target for Current School Year: Less than 3% of schools have >10% of kindergarteners either conditional or overdue.

Area 8. Influenza

Goal 8.1: Strengthen capacity to protect against seasonal influenza and to prepare for a	
pandemic.	
Required Activities	Performance Measures
Goal 8.1 Activity a: To assist your LHD emergency preparedness lead in fulfilling its emergency preparedness grant requirements, utilize IZB-supplied influenza vaccine or other 317-funded ⁷ vaccines to support at least one mass immunization exercise/year. Confirm your LHD emergency preparedness program has entered all doses into CAIR within 14 days of administration, as per the emergency preparedness grant requirement. Note: PHEP exercises can be used to fulfill this requirement.	 Mass vaccination exercise completed by local health department, including immunization and preparedness program staff. Number of exercises as SLVEs reported via SLVE Survey.
Goal 8.1 Activity b: Utilize IZB-supplied	Number of doses of influenza vaccine
influenza vaccine to immunize jurisdiction	administered.

⁶ "Targeted schools are schools with 10 or more kindergarteners that reported greater than 10% of students conditionally admitted and/or overdue for required immunization.

⁷ If the LHD would like to use Pan Flu Funding or other emergency preparedness funding for vaccine purchase, please reach out to the Immunization Branch with your request.

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against influenza; doses may be shared with	Target # of doses must be at least 90% of
local partners.	previous season's total doses.

Glossary of Acronyms and Terms

Abbreviation or term	Definition
317 vaccine	Vaccine provided to LHD clinics and partners for uninsured adults and for outbreak purposes.
ACIP	Advisory Committee on Immunization Practices
ACOG	American College of Obstetricians and Gynecologists
AIAW	Adolescent Immunization Action Week
CAIR	California Immunization Registry
CalREDIE	California Reportable Disease Information Exchange
CDC	Centers for Disease Control and Prevention
CDPH	California Department of Public Health
CIC	California Immunization Coalition
COVID-19	Coronavirus Disease 2019
CPSP	Comprehensive Perinatal Services Program
CSTE	Council of State and Territorial Epidemiologists
DNA	Deoxyribonucleic Acid
eCR	Electronic Case Reporting
EHR	Electronic Health Record
ELR	Electronic Laboratory Reporting
EZIZ	An Immunization Branch-operated website (eziz.org) with immunization training and resource materials.
FQHC	Federally Qualified Health Center
HBsAg	Hepatitis B Surface Antigen
HBV	Hepatitis B Vaccine
IgM	Immunoglobulin
IHS	Indian Health Services

Immunization Information System		
Immunization Quality Improvement for Providers		
Immunization Skills Institute		
Immunization		
Immunization Branch (of CDPH)		
Vaccine ordered through the CDPH Immunization Branch and supplied to LHD clinics or partners using state (SGF) or federal (VFC and 317) funding sources.		
Local CAIR representative (on CDPH IZB staff)		
Local Health Department		
Clinic run or housed in LHD that serves as a medical home for its patients. Includes federally qualified health centers or look-alikes that are operated or housed in LHDs		
Local Health Jurisdiction		
Medical Assistant		
Maternal Child and Adolescent Health		
Medi-Cal Managed Care Plan		
Memorandum of Understanding		
National Immunization Awareness Month		
National Influenza Vaccine Week		
Polymerase Chain Reaction		
Post Exposure Prophylaxis		
Perinatal Hepatitis B Prevention Program		
Provider Identification Number		
Post-Vaccination Serology		
School and Childcare Roster Lookup		
State General Fund		
School-Located Vaccine Event		
Tetanus, Diphtheria, and Pertussis		
Transitional Kindergarten/Kindergarten		

VFC	Vaccines for Children Program		
VPDs	Vaccine-Preventable Disease(s)		
VRDL	Viral and Rickettsial Disease Laboratory (of CDPH)		
WIC	Women, Infants, and Children		

AWARD ATTACHMENTS

California Department of Public Health Grant Award #: 1 NH23IP922690-01-00

Terms and Conditions

U.S. DEPARTMENT OF HEALTH AND HUMAN SERVICES

Public Health Service

Centers for Disease Control and Prevention (CDC) Atlanta GA 30333

General Terms and Conditions for Non-Research Grants and Cooperative Agreements

Incorporation: The U.S. Department of Health and Human Services (HHS) grant recipients must comply with: all terms and conditions outlined in the Notice of Funding Opportunity (NOFO); their Notice of Award (NOA); grants policy contained in applicable HHS Grants Policy Statements; HHS grant administration regulations (e.g., 45 CFR Part 75, 2 CFR 200 (as applicable)); requirements imposed by program statutes and regulations; applicable Executive Orders; HHS Administrative and National Policy Requirements; HHS policies, directives, and guidance; and requirements or limitations in any applicable appropriations acts. The term grant is used throughout these general terms and conditions of award and includes cooperative agreements.

Note: In the event that any requirement in the NOA, the NOFO, the HHS Grants Policy Statement, 45 CFR Part 75, or applicable statutes/appropriations acts conflict, then statutes and regulations take precedence.

Applicability of 2 CFR 200 Provisions Beginning October 1, 2024

This award is subject to the requirements in 45 CFR Part 75, except as amended by the following provisions of 2 CFR Part 200, which apply to new, continuation, and supplemental awards made on or after October 1, 2024.

- 2 CFR § 200.1. Definitions, "Modified Total Direct Cost", "Equipment", and "Supplies"
- 2 CFR § 200.313(e). Equipment, Disposition
- 2 CFR § 200.314(a). Supplies
- 2 CFR § 200.320. Procurement methods
- 2 CFR § 200.333. Fixed amount subawards
- 2 CFR § 200.344. Closeout
- 2 CFR § 200.414(f). Indirect costs, De Minimis Rate
- 2 CFR § 200.501. Audit requirements

2 CFR 200 citation	Replaces 45 CFR 75 citation		
2 CFR § 200.1. Definitions, "Modified Total	45 CFR § 75.2. Definitions, "Modified Total		
Direct Cost"	Direct Cost"		
2 CFR § 200.1. Definitions, "Equipment"	45 CFR § 75.2. Definitions, "Equipment"		
2 CFR § 200.1. Definitions, "Supplies"	45 CFR § 75.2. Definitions, "Supplies"		
2 CFR § 200.313(e). Equipment, Disposition	45 CFR § 75.320(e). Equipment, Disposition		
2 CFR § 200.314(a). Supplies	45 CFR § 75.321(a). Supplies		

2 CFR § 200.320. Procurement methods	45 CFR § 75.329. Procurement procedures	
2 CFR § 200.333. Fixed amount subawards	45 CFR § 75.353. Fixed amount subawards	
2 CFR § 200.344. Closeout	45 CFR § 75.381. Closeout	
2 CFR § 200.414(f). Indirect costs, De	45 CFR § 75.414(f). Indirect (F&A) costs, De	
Minimis Rate	Minimis Rate	
2 CFR § 200.501. Audit requirements	45 CFR § 75.501. Audit requirements	

FEDERAL REGULATIONS AND POLICIES

2 CFR 200 – Uniform Administrative Requirements, Cost Principles, and Audit Requirements for Federal Awards. Referenced where indicated and applicable.

https://www.ecfr.gov/current/title-2/subtitle-A/chapter-II/part-200?toc=1

45 CFR Part 75 – Uniform Administrative Requirements, Cost Principles, and Audit Requirements for HHS Awards. https://www.ecfr.gov/cgi-bin/text-idx?node=pt45.1.75&rgn=div5

HHS Administrative and National Policy Requirements

https://www.hhs.gov/sites/default/files/hhs-administrative-national-policy-requirements.pdf

HHS Grants Policy and Regulations

https://www.hhs.gov/grants/grants/grants-policies- regulations/index.html

HHS Grants Policy Statement (effective for new, continuation, and supplemental awards made on or after October 1, 2024)

https://www.hhs.gov/sites/default/files/hhs-grants-policy-statement-october-2024.pdf

HHS Grants Policy Statement (January 2007 version applies to awards issued before October 1, 2024)

https://public3.pagefreezer.com/browse/HHS.gov/27-09-

2024T06:59/https://www.hhs.gov/sites/default/files/grants/grants/policies-regulations/hhsgps107.pdf

Federal Funding Accountability and Transparency Act (FFATA). https://sam.gov/fsrs.

Refer to the section below on Reporting Requirements for more details.

Trafficking In Persons: Consistent with 2 CFR 175, awards are subject to the requirements of the Trafficking Victims Protection Act of 2000, as amended (22 U.S.C. Part 7104(g)). https://www.ecfr.gov/current/title-2/subtitle-A/chapter-I/part-175

FUNDING RESTRICTIONS AND LIMITATIONS

Cost Limitations as stated in Appropriations Acts. Recipients must follow applicable fiscal year appropriations law in effect at the time of award and consistent with the specific funds provided under that award. The general provisions for grants, cooperative agreements and loans funded by the Departments of Labor, Health and Human Services, Education, and Related Agencies Appropriations Act is available

at: https://www.congress.gov/resources/display/content/Appropriations+and+Budget.

Though Recipients are required to comply with all applicable appropriations restrictions, please

find below specific ones of note. CDC notes that the cited section for each below provision may change annually.

- A. Cap on Salaries (Division H, Title II, General Provisions, Sec. 202): None of the funds appropriated in this title shall be used to pay the salary of an individual, through a grant or other extramural mechanism, at a rate in excess of Executive Level II.
 - Note: The salary rate limitation does not restrict the salary that an organization may pay an individual working under an HHS award or order; it merely limits the portion of that salary that may be paid with federal funds. The HHS Grants Policy Statement further explains the application of this salary rate limitation.
- B. Gun Control Prohibition (Div. H, Title II, Sec. 210): None of the funds made available in this title may be used, in whole or in part, to advocate or promote gun control.
- C. Lobbying Restrictions (Div. H, Title V, Sec. 503):
 - 503(a): No part of any appropriation contained in this Act or transferred pursuant to section 4002 of Public Law 111-148 shall be used, other than for normal and recognized executive- legislative relationships, for publicity or propaganda purposes, for the preparation, distribution, or use of any kit, pamphlet, booklet, publication, electronic communication, radio, television, or video presentation designed to support or defeat the enactment of legislation before the Congress or any State or local legislature or legislative body, except in presentation to the Congress or any State or local legislature itself, or designed to support or defeat any proposed or pending regulation, administrative action, or order issued by the executive branch of any State or local government itself.
 - 503(b): No part of any appropriation contained in this Act or transferred pursuant to section 4002 of Public Law 111-148 shall be used to pay the salary or expenses of any grant or contract recipient, or agent acting for such recipient, related to any activity designed to influence the enactment of legislation, appropriations, regulation, administrative action, or Executive order proposed or pending before the Congress or any State government, State legislature or local legislature or legislative body, other than for normal and recognized executive-legislative relationships or participation by an agency or officer of a State, local or tribal government in policymaking and administrative processes within the executive branch of that government.
 - 503(c): The prohibitions in subsections (a) and (b) shall include any activity to advocate
 or promote any proposed, pending or future federal, state or local tax increase, or any
 proposed, pending, or future requirement or restriction on any legal consumer product,
 including its sale of marketing, including but not limited to the advocacy or promotion of
 gun control.

For additional information, see Anti-Lobbying Restrictions for CDC Grantees at https://www.cdc.gov/grants/documents/Anti-Lobbying-Restrictions.pdf.

D. Blocking access to pornography (Div. H, Title V, Sec. 520): (a) None of the funds made available in this Act may be used to maintain or establish a computer network unless such

network blocks the viewing, downloading, and exchanging of pornography; (b) Nothing in subsection (a) shall limit the use of funds necessary for any federal, state, tribal, or local law enforcement agency or any other entity carrying out criminal investigations, prosecution, or adjudication activities.

E. Needle Exchange (Div. H, Title V, Sec. 526): Notwithstanding any other provision of this Act, no funds appropriated in this Act shall be used to carry out any program of distributing sterile needles or syringes for the hypodermic injection of any illegal drug.

Prohibition on certain telecommunications and video surveillance services or equipment (2 CFR 200.216): For all new, non-competing continuation, renewal or supplemental awards issued on or after August 13, 2020, recipients and subrecipients are prohibited from obligating or expending grant funds (to include direct and indirect expenditures as well as cost share and program funds) to:

- 1. Procure or obtain,
- 2. Extend or renew a contract to procure or obtain; or
- 3. Enter into contract (or extend or renew contract) to procure or obtain equipment, services, or systems that use covered telecommunications equipment or services as a substantial or essential component of any system, or as critical technology as part of any system. As described in <u>2 CFR 200.216</u>, covered telecommunications equipment is telecommunications equipment produced by Huawei Technologies Company or ZTE Corporation (or any subsidiary or affiliate of such entities).
 - i. For the purpose of public safety, security of government facilities, physical security surveillance of critical infrastructure, and other national security purposes, video surveillance and telecommunications equipment produced by Hytera Communications Corporation, Hangzhou Hikvision Digital Technology Company, or Dahua Technology Company (or any subsidiary or affiliate of such entities).
 - ii. Telecommunications or video surveillance services provided by such entities or using such equipment.
 - iii. Telecommunications or video surveillance equipment or services produced or provided by an entity that the Secretary of Defense, in consultation with the Director of the National Intelligence or the Director of the Federal Bureau of Investigation, reasonably believes to be an entity owned or controlled by, or otherwise, connected to the government of a covered foreign country.

President's Emergency Plan for AIDS Relief (PEPFAR) funding is exempt from the prohibition under <u>2 CFR 200.216</u> until September 30, 2028. During the exemption period, PEPFAR recipients are expected to work toward implementation of <u>2 CFR 200.216</u>. The exemption may only be applied when there is no available alternative eligible source for these services.

Cancel Year: 31 U.S.C. Part 1552(a) Procedure for Appropriation Accounts Available for Definite Periods states the following: On September 30th of the 5th fiscal year after the period of availability for obligation of a fixed appropriation account ends, the account shall be closed and any remaining balances (whether obligated or unobligated) in the account shall be canceled and thereafter shall not be available for obligation or expenditure for any purpose.

REPORTING REQUIREMENTS

Annual Federal Financial Report (FFR, SF-425): The Annual Federal Financial Report (FFR) SF-425 is required and must be submitted no later than 90 days after the end of the budget period in the Payment Management System.

Additional guidance on submission of Federal Financial Reports can be found at https://www.cdc.gov/grants/documents/change-in-federal-reporting-fy-2021-recipients.pdf.

If more frequent reporting is required, the Notice of Award terms and conditions will explicitly state the reporting requirement.

Annual Performance Progress and Monitoring Reporting: The Annual Performance Progress and Monitoring Report (PPMR) is due no later than 120 days prior to the end of the budget period and serves as the continuation application for the follow-on budget period. Submission instructions, due date, and format will be included in the guidance from the assigned GMO/GMS via www.grantsolutions.gov.

Any change to the existing information collection noted in the award terms and conditions will be subject to review and approval by the Office of Management and Budget (OMB) under the Paperwork Reduction Act.

Data Collection and Sharing Under Award: Consistent with strategies and activities expected and anticipated under this award, Recipient, either directly or indirectly, may be expected to collect or generate data for public health purposes. For purposes of this award, data for public health purposes may be administrative data or data commonly accepted in the scientific community as a basis for public health findings, conclusions, and implementation, but does not include preliminary analyses, drafts of scientific papers, plans for future research communications with colleagues, or physical objects, such as laboratory notebooks or laboratory specimens unless otherwise specified in the award.

45 C.F.R. 75.322(d) states that the federal government has the right to: 1) obtain, reproduce, publish, or otherwise use the data produced under a <u>federal award</u>; and 2) authorize others to receive, reproduce, publish, or otherwise use such data for federal purposes. In furtherance of various United States Government-wide initiatives and policies, the federal government seeks to make federally funded publications and data underlying them more readily available, and to make public health data more readily accessible within the federal government and to the public. Consistent with grant regulations, CDC may legally obtain a copy of any data collected or generated under this award. Where CDC has determined that data collected or generated under this award must be shared with CDC, such direction will be further addressed in your Notice of Funding Opportunity, your Notice of Grant Award, or other specific grant guidance. Acceptance of funds under this award is an acknowledgement of this regulatory provision and its application to this award.

Data Management Plan: CDC requires recipients for projects that involve the collection or generation of data with federal funds to develop, submit, and comply with a Data Management Plan (DMP) for each collection or generation of public health data undertaken as part of the award. The DMP should take into consideration sharing data with CDC including: 1) the specific data that will be shared under the award, 2) the process and timing planned for such sharing, 3) and any

legal limitations that the Recipient asserts would hinder CDC access to, or use of, the data collected or generated under the award. In addition, the DMP should address broader access to and archiving/long-term preservation of collected or generated data. Additional information on the Data Management and Access requirements can be found at https://www.cdc.gov/grants/additional-requirements/ar-25.html.

Audit Requirement Domestic Organizations (including US-based organizations implementing projects with foreign components): An organization that expends \$1,000,000 or more in a fiscal year in federal awards shall have a single or program-specific audit conducted for that year in accordance with the provisions of 2 CFR 200.501. The audit period is an organization's fiscal year. The audit must be completed along with a data collection form (SF-SAC), and the reporting package shall be submitted within the earlier of 30 days after receipt of the auditor's report(s), or nine (9) months after the end of the audit period. The audit report must be sent to:

Federal Audit Clearing House Internet Data Entry System Electronic Submission: https://harvester.census.gov/facides/(S(0vkw1zaelyzjibnahocga5i0))/account/login.aspx

AND

Office of Financial Resources, Office of Risk Management and Internal Controls, Audit Resolution Team (ART), ORMIC.Audit.Resolution@cdc.gov.

Audit Requirement Foreign Organizations: A foreign organization that expends \$300,000 or more in a fiscal year on its federal awards must have a single or program-specific audit conducted for that year. The audit period is an organization's fiscal year. The auditor shall be a U.S.-based Certified Public Accountant firm, the foreign government's Supreme Audit Institution or equivalent, or an audit firm endorsed by the U.S. Agency for International Development's Office of Inspector General. The audit must be completed in English and in US dollars and submitted within the earlier of 30 days after receipt of the auditor's report(s), or nine (9) months after the end of the audit period. The audit report must be sent to the Office of Financial Resources, Office of Risk Management and Internal Controls, Audit Resolution Team (ART) at ORMIC.Audit.Resolution@cdc.gov. After receipt of the audit report, CDC will resolve findings by issuing Final Management Determination Letters.

Domestic and Foreign organizations: Audit requirements for Subrecipients to whom 45 CFR 75 Subpart F applies: The recipient must ensure that the subrecipients receiving CDC funds also meet these requirements. The recipient must also ensure to take appropriate corrective action within six months after receipt of the subrecipient audit report in instances of non-compliance with applicable federal law and regulations (45 CFR 75 Subpart F and HHS Grants Policy Statement). The recipient may consider whether subrecipient audits necessitate adjustment of the recipient's own accounting records. If a subrecipient is not required to have a program-specific audit, the recipient is still required to perform adequate monitoring of subrecipient activities. The recipient shall require each subrecipient to permit the independent auditor access to the subrecipient's records and financial statements. The recipient must include this requirement in all subrecipient contracts.

Federal Funding Accountability and Transparency Act (FFATA)

In accordance with 2 CFR Chapter 1, Part 170 Reporting Sub-Award and Executive Compensation Information, Prime Recipients awarded a federal grant are required to file a FFATA sub-award report

by the end of the month following the month in which the prime recipient awards any sub-grant equal to or greater than \$30,000. Refer to 2 CFR Chapter 1, Part 170 Reporting Sub-Award and Executive Compensation Information at <u>eCFR :: 2 CFR Part 170 -- Reporting Subaward and Executive Compensation Information and https://exam.gov/fsrs for reporting requirements and guidance.</u>

Unique Entity Identifier (UEI)

The UEI is the official identifier for doing business with the U.S. Government as of April 4, 2022. The UEI is generated and assigned by the System for Award Management at SAM.gov. In accordance with 2 CFR part 25, Appendix A, a recipient must maintain current information in SAM.gov, through at least annual review, until it submits the final required financial report or receives the final payment, whichever is later.

Required Disclosures for Responsibility and Qualification (R/Q) (SAM.gov): Consistent with 45 CFR 75.113, applicants and recipients must disclose in a timely manner, in writing to the CDC, with a copy to the HHS Office of Inspector General (OIG), all information related to violations of federal criminal law involving fraud, bribery, or gratuity violations potentially affecting the federal award. Subrecipients must disclose, in a timely manner in writing to the prime recipient (pass through entity) and the HHS OIG, all information related to violations of federal criminal law involving fraud, bribery, or gratuity violations potentially affecting the federal award. Disclosures must be sent in writing to the assigned GMS/GMO identified in the NOA, and to the HHS OIG by email at grantdisclosures@oig.hhs.gov or by mail to the following address:

U.S. Department of Health and Human Services
Office of the Inspector General
ATTN: Mandatory Grant Disclosures, Intake Coordinator
330 Independence Avenue, SW
Cohen Building, Room 5527
Washington, DC 20201

Recipients must include this mandatory disclosure requirement in all subawards and contracts under this award.

Failure to make required disclosures can result in any of the remedies described in 45 CFR 75.371. Remedies for noncompliance include suspension or debarment (See 2 CFR parts 180 and 376, and 31 U.S.C. 3321).

CDC is required to report any termination of a federal award prior to the end of the period of performance due to material failure to comply with the terms and conditions of this award in the OMB-designated Responsibilities and Qualifications (R/Q) accessible through SAM (45 CFR 75.372(b)). CDC must also notify the recipient if the federal award is terminated for failure to comply with the federal statutes, regulations, or terms and conditions of the federal award (45 CFR 75.373(b)).

1. General Reporting Requirement

If the total value of currently active grants, cooperative agreements, and procurement contracts from all federal awarding agencies exceeds \$10,000,000 for any period of time during the period of performance of this federal award, the recipient must maintain the currency of information reported to the System for Award Management (SAM) and made

available in the designated integrity and performance system (currently the Responsibility/Qualification (R/Q) through SAM.gov) about civil, criminal, or administrative proceedings described in section 2 of this award term and condition. This is a statutory requirement under section 872 of Public Law 110-417, as amended (41 U.S.C. 2313). As required by section 3010 of Public Law 111-212, all information posted in the designated integrity and performance system on or after April 15, 2011, except past performance reviews required for federal procurement contracts, will be publicly available.

2. Proceedings About Which You Must Report

Submit the information required about each proceeding that:

- a. Is in connection with the award or performance of a grant, cooperative agreement, or procurement contract from the federal government;
- b. Reached its final disposition during the most recent five-year period; and
- c. If one of the following:
 - (1) A criminal proceeding that resulted in a conviction, as defined in paragraph 5 of this award term and condition;
 - (2) A civil proceeding that resulted in a finding of fault and liability and payment of a monetary fine, penalty, reimbursement, restitution, or damages of \$5,000 or more;
 - (3) An administrative proceeding, as defined in paragraph 5 of this award term and condition, that resulted in a finding of fault and liability and your payment of either a monetary fine or penalty of \$5,000 or more or reimbursement, restitution, or damages in excess of \$100,000; or
 - (4) Any other criminal, civil, or administrative proceeding if:
 - (i) It could have led to an outcome described in paragraph 2.c.(1), (2), or (3) of this award term and condition;
 - (ii) It had a different disposition arrived at by consent or compromise with an acknowledgement of fault on your part; and
 - (iii) The requirement in this award term and condition to disclose information about the proceeding does not conflict with applicable laws and regulations.

3. Reporting Procedures

Enter in the SAM Entity Management area the information that SAM requires about each proceeding described in section 2 of this award term and condition. You do not need to submit the information a second time under assistance awards that you received if you already provided the information through SAM because you were required to do so under federal procurement contracts that you were awarded.

4. Reporting Frequency

During any period of time when you are subject to this requirement in section 1 of this award term and condition, you must report proceedings information through SAM for the most recent five-year period, either to report new information about any proceeding(s) that you

have not reported previously or affirm that there is no new information to report. Recipients that have federal contract, grant, and cooperative agreement awards with a cumulative total value greater than \$10,000,000 must disclose semiannually any information about the criminal, civil, and administrative proceedings.

5. Definitions

For purposes of this award term and condition:

- a. Administrative proceeding means a non-judicial process that is adjudicatory in nature in order to make a determination of fault or liability (e.g., Securities and Exchange Commission Administrative proceedings, Civilian Board of Contract Appeals proceedings, and Armed Services Board of Contract Appeals proceedings). This includes proceedings at the federal and state level but only in connection with performance of a federal contract or grant. It does not include audits, site visits, corrective plans, or inspection of deliverables.
- b. Conviction, for purposes of this award term and condition, means a judgment or conviction of a criminal offense by any court of competent jurisdiction, whether entered upon a verdict or a plea, and includes a conviction entered upon a plea of nolo contendere.
- c. Total value of currently active grants, cooperative agreements, and procurement contracts includes—
 - (1) Only the federal share of the funding under any federal award with a recipient cost share or match;
 - (2) The value of all expected funding increments under a federal award and options, even if not yet exercised.

GENERAL REQUIREMENTS

You will administer your project in compliance with the HHS Administrative and National Policy Requirements found at https://www.hhs.gov/sites/default/files/hhs-administrative-national-policy-requirements.pdf.

Termination (45 CFR Part 75.372) applies to this award and states, in part, the following:

This award may be terminated in whole or in part:

- (1) By the HHS awarding agency or pass-through entity, if a non-Federal entity fails to comply with the terms and conditions of a Federal award:
- (2) By the HHS awarding agency or pass-through entity for cause;
- (3) By the HHS awarding agency or pass-through entity with the consent of the non-Federal entity, in which case the two parties must agree upon the termination conditions, including the effective date and, in the case of partial termination, the portion to be terminated;
- (4) By the non-Federal entity upon sending to the HHS awarding agency or pass-through entity written notification setting forth the reasons for such termination, the effective date, and, in the case of partial termination, the portion to be terminated.

However, if the Federal awarding agency or pass-through entity determines in the case of partial termination that the reduced or modified portion of the Federal award or subaward will not accomplish the purposes for which the Federal award was made, the HHS awarding agency or pass-through entity may terminate the Federal award in its entirety.

Travel Cost: In accordance with HHS Grants Policy Statement, travel costs are allowable when the travel will provide a direct benefit to the project or program. To prevent disallowance of cost, the recipient is responsible for ensuring travel costs are clearly stated in their budget narrative and are applied in accordance with their organization's established travel policies and procedures. The recipient's established travel policies and procedures must also meet the requirements of 45 CFR Part 75.474.

Food and Meals: Costs associated with food or meals are allowable when consistent with applicable federal regulations and HHS policies. See https://www.hhs.gov/grants/contracts/contract-policies-regulations/spending-on-food/index.html. In addition, costs must be clearly stated in the budget narrative and be consistent with organization approved policies. Recipients must make a determination of reasonableness and organization approved policies must meet the requirements of 45 CFR Part 75.432.

Prior Approval: All requests which require prior approval, must bear the signature (or electronic authorization) of the authorized organization representative. The recipient should submit these requests no later than 120 days prior to the budget period's end date to ensure ample time remains to process and carry-out the request. Additionally, any requests involving funding issues must include an itemized budget and a narrative justification of the request.

The following types of requests are examples of actions that require prior approval, unless an expanded authority, or conversely a high-risk condition, is explicitly indicated in the NOA.

- Use of unobligated funds from prior budget period (Carryover)
- Lift funding restriction
- Significant redirection of funds (i.e., cumulative changes of 25% of total award)
- Change in scope
- Implement a new activity or enter into a sub-award that is not specified in the approved budget
- Apply for supplemental funds
- Extensions to period of performance

Templates for prior approval requests can be found at: https://www.cdc.gov/grants/already-have-grant/prior-approval-requests.html.

Additional information on the electronic grants administration system CDC non-research awards utilize, GrantSolutions, can be found at: https://www.cdc.gov/grants/grantsolutions/index.html.

Recipient Contractual/Consultant Cost Agreements: In accordance with §2 CFR 200.325, all supporting documentation related to the elements outlined in the <u>Budget Preparation Guidelines</u> must be maintained by the recipient and available upon request. Recipients may submit supporting documentation via GrantSolutions Grants Management Services (GSGMS) Grant Notes to the

assigned Grants Management Specialist.

Key Personnel: In accordance with 45 CFR Part 75.308, CDC recipients must obtain prior approval from CDC for (1) change in the project director/principal investigator, authorized organizational representative, business official, financial director, or other key persons specified in the NOFO, application or award document; and (2) the disengagement from the project for more than three months, or a 25 percent reduction in time devoted to the project, by the approved project director or principal investigator.

Inventions: Acceptance of grant funds obligates recipients to comply with the standard patent rights clause in 37 CFR Part 401.14.

Acknowledgment of Federal Funding: When issuing statements, press releases, publications, requests for proposal, bid solicitations and other documents --such as toolkits, resource guides, websites, and presentations (hereafter "statements") --describing the projects or programs funded in whole or in part with U.S. Department of Health and Human Services (HHS) federal funds, the recipient must clearly state:

- 1. the percentage and dollar amount of the total costs of the program or project funded with federal money; and,
- 2. the percentage and dollar amount of the total costs of the project or program funded by non-governmental sources.

When issuing statements resulting from activities supported by HHS financial assistance, the recipient entity must include an acknowledgement of federal assistance using one of the following or a similar statement.

If the HHS Grant or Cooperative Agreement is <u>NOT</u> funded with other non-governmental sources:

This [project/publication/program/website, etc.] [is/was] supported by the Centers for Disease Control and Prevention of the U.S. Department of Health and Human Services (HHS) as part of a financial assistance award totaling **\$XX** with 100 percent funded by CDC/HHS. The contents are those of the author(s) and do not necessarily represent the official views of, nor an endorsement, by CDC/HHS, or the U.S. Government.

If the HHS Grant or Cooperative Agreement IS partially funded with other non-governmental sources:

This [project/publication/program/website, etc.] [is/was] supported by the Centers for Disease Control and Prevention of the U.S. Department of Health and Human Services (HHS) as part of a financial assistance award totaling \$XX with XX percentage funded by CDC/HHS and \$XX amount and XX percentage funded by non- government source(s). The contents are those of the author(s) and do not necessarily represent the official views of, nor an endorsement, by CDC/HHS, or the U.S. Government.

The federal award total must reflect total costs (direct and indirect) for all authorized funds (including supplements and carryover) for the total competitive segment up to the time of the public statement.

Any amendments by the recipient to the acknowledgement statement must be coordinated with the HHS Awarding Agency.

If the recipient plans to issue a press release concerning the outcome of activities supported by HHS financial assistance, it should notify the HHS Awarding Agency in advance to allow for coordination.

Copyright Interests Provision: This provision is intended to ensure that the public has access to the results and accomplishments of public health activities funded by CDC. Pursuant to applicable grant regulations and CDC's Public Access Policy, Recipient agrees to submit into the National Institutes of Health (NIH) Manuscript Submission (NIHMS) system an electronic version of the final, peer-reviewed manuscript of any such work developed under this award upon acceptance for publication, to be made publicly available without any embargo or delay after publication. Also, at the time of submission, Recipient and/or the Recipient's submitting author must also post the manuscript through PubMed Central (PMC) without any embargo or delay after publication. The recipient must obtain prior approval from the CDC for any exception to this provision.

The author's final, peer-reviewed manuscript is defined as the final version accepted for journal publication and includes all modifications from the publishing peer review process, and all graphics and supplemental material associated with the article. Recipient and its submitting authors working under this award are responsible for ensuring that any publishing or copyright agreements concerning submitted article reserve adequate right to fully comply with this provision and the license reserved by CDC.

The manuscript will be hosted in both PMC and the CDC Stacks institutional repository system. In progress reports for this award, recipient must identify publications subject to the CDC Public Access Policy by using the applicable NIHMS identification number for up to three (3) months after the publication date and the PubMed Central identification number (PMCID) thereafter.

Disclaimer for Conference/Meeting/Seminar Materials: If a conference/meeting/seminar is funded by a grant, cooperative agreement, sub-grant and/or a contract, the recipient must include the following statement on conference materials, including promotional materials, agenda, and internet sites:

Funding for this conference was made possible (in part) by the Centers for Disease Control and Prevention. The views expressed in written conference materials or publications and by speakers and moderators do not necessarily reflect the official policies of the Department of Health and Human Services, nor does the mention of trade names, commercial practices, or organizations imply endorsement by the U.S. Government.

Logo Use for Conference and Other Materials: Neither the Department of Health and Human Services (HHS) nor the CDC logo may be displayed if such display would cause confusion as to the funding source or give false appearance of Government endorsement. Use of the HHS name or logo is governed by U.S.C. Part 1320b-10, which prohibits misuse of the HHS name and emblem in written communication. As a general matter, a non-federal entity is not authorized to use the HHS name or logo. Moreover, the HHS Office of the Inspector General has authority to

impose civil monetary penalties for violations (42 CFR Part 1003). The appropriate use of the HHS logo is subject to review and approval of the HHS Assistant Secretary for Public Affairs (ASPA), and if granted would be governed by a logo license agreement setting forth the terms and conditions of use.

Additionally, the CDC logo cannot be used by the recipient without the express, written consent of CDC, generally in the form of a logo license agreement setting forth the terms and conditions of use. The Program Official/Project Officer identified in the NOA can assist with facilitating such a request. It is the responsibility of the recipient to request consent for use of the logo in sufficient detail to ensure a complete depiction and disclosure of all uses of the Government logos. In all cases for utilization of Government logos, the recipient must ensure written consent is received.

Equipment and Products: To the greatest extent practical, all equipment and products purchased with CDC funds should be American made. CDC defines equipment as tangible non-expendable personal property (including exempt property) charged directly to an award having a useful life of more than one year AND an acquisition cost of \$10,000 or more per unit. However, consistent with recipient policy, a lower threshold may be established. Please provide the information to the Grants Management Officer to establish a lower equipment threshold to reflect your organization's policy. The recipient may use its own property management standards and procedures, provided it observes provisions in applicable grant regulations found at 45 CFR Part 75.

Federal Information Security Management Act (FISMA): All information systems, electronic or hard copy, that contain federal data must be protected from unauthorized access. This standard also applies to information associated with CDC grants. Congress and the OMB have instituted laws, policies and directives that govern the creation and implementation of federal information security practices that pertain specifically to grants and contracts. The current regulations are pursuant to the Federal Information Security Management Act (FISMA), Title III of the E-Government Act of 2002, PL 107-347.

FISMA applies to CDC recipients only when recipients collect, store, process, transmit or use information on behalf of HHS or any of its component organizations. In all other cases, FISMA is not applicable to recipients of grants, including cooperative agreements. Under FISMA, the recipient retains the original data and intellectual property, and is responsible for the security of these data, subject to all applicable laws protecting security, privacy, and research. If/When information collected by a recipient is provided to HHS, responsibility for the protection of the HHS copy of the information is transferred to HHS and it becomes the agency's responsibility to protect that information and any derivative copies as required by FISMA.

For the full text of the requirements under Federal Information Security Management Act (FISMA), Title III of the E- Government Act of 2002 Pub. L. No. 107-347, please review the following website: https://www.govinfo.gov/content/pkg/PLAW-107publ347/pdf/PLAW-107publ347.pdf.

Whistleblower Protections: As a recipient of this award, you must comply with the National Defense Authorization Act (NDAA) for Fiscal Year (FY) 2013 (Pub. L. 112-239, 41 U.S.C. § 4712) "Enhancement of contractor protection from reprisal for disclosure of certain information," and 48 CFR part 3 subpart 3.9, "Whistleblower Protections for Contractor Employees." For more information see:

https://oig.hhs.gov/fraud/whistleblower/.

Cybersecurity Requirements: Recipients shall develop plans and procedures, modeled after the NIST Cybersecurity framework, to protect HHS and CDC systems and data, if the following conditions are met: 1) recipients, subrecipients, or third-party entities have ongoing and consistent access to HHS owned or operated information or operational technology systems and 2) recipients, subrecipients, or third-party entities receive, maintain, transmit, store, access, exchange, process, or utilize personal identifiable information (PII) or personal health information (PHI) obtained from the awarding HHS agency for the purposes of executing the award. Where both conditions exist, recipients must develop cybersecurity plans and procedures modeled after the NIST Cybersecurity framework (https://www.nist.gov/cyberframework) to protect HHS systems and data.

PAYMENT INFORMATION

Fraud Waste or Abuse: The HHS Office of the Inspector General (OIG) maintains a toll-free number (1-800-HHS-TIPS [1-800-447-8477]) for receiving information concerning fraud, waste, or abuse under grants and cooperative agreements. Information also may be submitted online at https://tips.oig.hhs.gov/ or by mail to U.S. Department of Health and Human Services, Office of the Inspector General, Attn: OIG HOTLINE OPERATIONS, P.O. Box 23489 Washington DC 20026. Such reports are treated as sensitive material and submitters may decline to give their names if they choose to remain anonymous.

For additional information, see: https://oig.hhs.gov/fraud/report-fraud/.

Automatic Drawdown (Direct/Advance Payments): Payments under CDC awards will be made available through the Department of Health and Human Services (HHS) Payment Management System (PMS), under automatic drawdown, unless specified otherwise in the NOA. Recipients must comply with requirements imposed by the PMS on-line system. Questions concerning award payments or audit inquiries should be directed to the payment management services office.

PMS Website: https://pms.psc.gov/
PMS Phone Support: +1(877)614-5533
PMS Email Support: PMSSupport@psc.gov

Payment Management System Subaccount: Funds awarded in support of approved activities will be obligated in an established subaccount in the PMS. Funds must be used in support of approved activities in the NOFO and the approved application. All award funds must be tracked and reported separately.

Exchange Rate: All requests for funds contained in the budget, shall be stated in U.S. dollars. Once an award is made, CDC will generally not compensate foreign recipients for currency exchange fluctuations through the issuance of supplemental awards.

Acceptance of the Terms of an Award: By drawing or otherwise obtaining funds from PMS, the recipient acknowledges acceptance of the terms and conditions of the award and is obligated to perform in accordance with the requirements of the award. If the recipient cannot accept the terms, the recipient should notify the Grants Management Officer within thirty (30) days of receipt of the NOA.

Certification Statement: By drawing down funds, the recipient certifies that proper financial management controls and accounting systems, to include personnel policies and procedures, have been established to adequately administer federal awards and funds drawn down. Recipients must comply with all terms and conditions in the NOFO, outlined in their NOA, grant policy terms and conditions contained in applicable HHS Grant Policy Statements, and requirements imposed by program statutes and regulations and HHS grants administration regulations, as applicable, as well as any regulations or limitations in any applicable appropriations acts.

CLOSEOUT REQUIREMENTS

In accordance with 2 CFR 200.344, recipients must submit all closeout reports identified in this section within 120 days of the period of performance end date. The reporting timeframe is the full period of performance. If the recipient does not submit all reports in accordance with this section and the terms and conditions of the Federal Award, CDC may proceed to close out with the information available within one year of the period of performance end date unless otherwise directed by authorizing statutes. Failure to submit timely and accurate final reports may affect future funding to the organization or awards under the direction of the same Project Director/Principal Investigator (PD/PI). If recipients do not submit all closeout reports identified in this section within one year of the period of performance end date, then CDC must report recipients' material failure to comply with the terms and conditions of the award with the OMB-designated integrity and performance system (currently Responsibility/Qualification section of SAM.gov). CDC may also pursue other enforcement actions per 45 CFR 75.371.

Final Performance Progress and Evaluation Report (PPER): This report should include the information specified in the NOFO and is submitted upon solicitation from the GMS/GMO via www.grantsolutions.gov. At a minimum, the report will include the following:

- Statement of progress made toward the achievement of originally stated aims;
- Description of results (positive or negative) considered significant; and
- List of publications resulting from the project, with plans, if any, for further publication.

All manuscripts published as a result of the work supported in part or whole by the grant must be submitted with the performance progress reports.

Final Federal Financial Report (FFR, SF-425): The FFR should only include those funds authorized and expended during the timeframe covered by the report. The Final FFR, SF-425 is required and must be submitted no later than 120 days after the period of performance end date through recipient online accounts in the Payment Management System. The final FFR will consolidate data reporting responsibilities to one entry point within PMS which will assist with the reconciliation of expenditures and disbursements to support the timely close-out of grants.

The final FFR must indicate the exact balance of unobligated funds and may not reflect any unliquidated obligations. Remaining unobligated funds will be de-obligated and returned to the U.S. Treasury.

Every recipient should already have a PMS account to allow access to complete the SF-425.

Additional guidance on submission of Federal Financial Reports can be found at https://www.cdc.gov/grants/documents/change-in-federal-reporting-fy-2021-recipients.pdf.

Equipment and Supplies - Tangible Personal Property Report (SF-428): A completed Tangible Personal Property Report SF-428 and Final Report SF-428B addendum must be submitted, along with any Supplemental Sheet SF-428S detailing all major equipment acquired or furnished under this project with a unit acquisition cost of \$10,000 or more. Electronic versions of the forms can be downloaded by visiting: https://www.grants.gov/forms/forms-repository/post-award-reporting-forms.

If no equipment was acquired under an award, a negative report is required. The recipient must identify each item of equipment that it wishes to retain for continued use in accordance with 45 CFR Part 75. The awarding agency may exercise its rights to require the transfer of equipment purchased under the assistance award. CDC will notify the recipient if transfer to title will be required and provide disposition instruction on all major equipment.

Equipment with a unit acquisition cost of less than \$10,000 that is no longer to be used in projects or programs currently or previously sponsored by the federal government may be retained, sold, or otherwise disposed of, with no further obligation to the federal government (see 2 CFR 200.313(e)(1)).

CDC STAFF RESPONSIBILITIES

Roles and Responsibilities: Grants Management Specialists/Officers (GMO/GMS) and Program Officials (PO) work together to award and manage CDC grants and cooperative agreements. From the pre-planning stage to closeout of an award, grants management and program staff have specific roles and responsibilities for each phase of the grant cycle. Award specific terms and conditions will include contact information for the PO/GMO/GMS.

Program Official: The PO is the federal official responsible for monitoring the programmatic, scientific, and/or technical aspects of grants and cooperative agreements including:

- The development of programs and NOFOs to meet the CDC's mission;
- Providing technical assistance to applicants in developing their applications, e.g., explanation of programmatic requirements, regulations, evaluation criteria, and guidance to applicants on possible linkages with other resources;
- Providing technical assistance to recipients in the performance of their project; and
- Post-award monitoring of recipient performance such as review of progress reports, review of prior approval requests, conducting site visits, and other activities complementary to those of the GMO/GMS.

For Cooperative Agreements, substantial involvement is required from CDC. The PO is the federal official responsible for the collaboration or participation in carrying out the effort under the award. Substantial involvement will be detailed in the NOFO and award specific terms and conditions and may include, but is not limited to:

- Review and approval of one stage of work before work can begin on a subsequent stage;
- Review and approval of substantive programmatic provisions of proposed subawards or contracts (beyond existing federal review of procurement or sole source policies);

- Involvement in the selection of key relevant personnel;
- CDC and recipient collaboration or joint participation; and
- Implementing highly prescriptive requirements prior to award limiting recipient discretion with respect to scope of services, organizational structure, staffing, mode of operation, and other management processes.

Grants Management Officer: The GMO is the only official authorized to obligate federal funds and is responsible for signing the NOA, including revisions to the NOA that change the terms and conditions. The GMO serves as the counterpart to the business officer of the recipient organization. The GMO is the federal official responsible for the business and other non-programmatic aspects of grant awards including:

- Determining the appropriate award instrument, i.e., grant or cooperative agreement;
- Determining if an application meets the requirements of the NOFO;
- Ensuring objective reviews are conducted in an above-the-board manner and according to guidelines set forth in grants policy;
- Ensuring recipient compliance with applicable laws, regulations, and policies;
- Negotiating awards, including budgets;
- Responding to recipient inquiries regarding the business and administrative aspects of an award;
- Providing recipients with guidance on the closeout process and administering the closeout of grants;
- Receiving and processing reports and prior approval requests such as changes in funding, budget redirection, or changes to the terms and conditions of an award; and
- Maintaining the official grant file and program book.

Grants Management Specialist: The GMS is the federal staff member responsible for the day-to-day management of grants and cooperative agreements. The GMS is the primary contact of recipients for business and administrative matters pertinent to grant awards. Many of the functions described in the GMO section are performed by the GMS, on behalf of the GMO.

Y2025-26 Federal Compliance Requirements of the Immunization Grant No.: 1 NH23IP922690-01-00							
This section requires Subrecipient signature to acknowledge that the Subrecipient has reviewed and understand the Federal Compliance Requirements of the Immunization Grant. See enclosed copy of the Award Attachments under which this funding is issued.							
Dawn Rowe, Chair, Board of Supervisors							
Print Name and Title of Person Signing Signature of Person Signing Date							

CDPH Immunization Branch Subrecipient Information Form Update FY 2025-26

Date Form Completed: 8/11/2025

	This is the information that will appear on your grant agreement cover page.					
	Federal Tax ID #	95-6002748	CDPH Contract/Grant#	(will be assigned by IZ/CDPH)		
	Unique Entity Identifier (UEI) #	PD18A8XKE7B6 Is this the County's	Department of Pub	lic Health UEI? X Yes □ No		
	Official Organization Name (Based on UEI)	San Bernardino County				
	Mailing Address	451 E. Vanderbilt Way, San Bernardino, CA 92415				
	Street Address (If Different)					
	County	San Bernardino	_			
	Phone	(800) 782-4264	_ Fax	(909) 381-8471		
on	Website	www.sbcounty.gov		-		
ati	County Personnel Salaries	⊠ Yes				
ganization	Available on the County Public Website	□ No				
Orga	Click this box if there were no changes from last year.	☐ No changes		•		