AMEN	NDMENT OF SOLICITATION	I/MODIFICATION	OF CONTRACT	1.0	JONIKACI ID C	ODE	PAGE 1 OF 4
2. AMENDMEI P00010	NT/MODIFICATION NO.	3. EFFECTIVE DATE See Blk. 16C	4. REQUISITION/PURG See Block 14	L CHASE	REQ. NO.	5. PROJECT	ΓΝΟ. (If applicable)
6. ISSUED BY	CODE	SPE300	7. ADMINISTERED BY (If	other t	han Item 6)	CODE	S4402A
700 ROBBINS A PHILADELPHIA USA Initiator: Steven	SUPPLY CHAIN AVENUE A PA 19111-5096		DCMA DALLAS 4211 CEDAR SPRIN DALLAS TX 75219-2 USA		OAD		
8. NAME AND	ADDRESS OF CONTRACTOR (No., street	t, county, State and ZIP	Code)	(X)	9A. AMENDMEN	NT OF SOLICITA	ATION NO.
Sterling Food Sterling Food 1075 Arion P SAN ANTON USA	s, LLC			X		TION OF CONT 17-D-Z109	RACT/ORDER NO.
CODE 7M71	2 FAG	CILITY CODE				2016 NOV (08
	11. THIS ITEM	ONLY APPLIES TO A	MENDMENTS OF SO	LICI	TATIONS		
(a) By completing or (c) By separa PLACE DESIGN amendment, and amendment, and amendment, and amendment.		copies of the amendment to the solicitation and amendr OR TO THE HOUR AND DATE such change may be made by to e specified. APPLIES ONLY TO MO ES THE CONTRACT/OI SUANT TO: (Specify author DER IS MODIFIED TO REFLEC UANT TO THE AUTHORITY OF SENTERED INTO PURSUAN	t; (b) By acknowledging receinent numbers. FAILURE OF SPECIFIED MAY RESULT I elegram or letter, provided each of the second of t	ONTF	is amendment on ACKNOWLEDGM ECTION OF YOUR gram or letter make RACTS/ORDE D IN ITEM 14 ARE IN ITEM 14 ARE I	each copy of the ENT TO BE REC R OFFER. If by v es reference to t ERS, MADE IN THE C	CEIVED AT THE irtue of this the solicitation and this contract or the solicitation and the solici
	ANT: Contractor X is not, Org	is required to sign this				es to the issu	
Except as prov 15A NAME ANI	ided herein, all terms and conditions of the do	ocument referenced in Item 9/	A or 10A, as heretofore chan 16A. NAME AND TITLE OI KATHERINE KNECHT 16B. UNITED STATES OF	F CON	TRACTING OFFI		
IJD. CONTRAC	OTOTYOTT LINON	.ss. bitte diditeb		L			2020 DEC 01
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(Signature of person authorized to sign)

(Signature of Contracting Officer)

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Part 12 Clauses

- I. Due to system issues, an older modification has overwritten the language for modification P00009; therefore modification P00009 is hereby deleted in its entirety and replaced with modification P00010
- II. The following changes apply to Contract SPE3S1-17-D-Z109 at no additional cost:
- A. Solicitation SPE3S1-16-R-0007, page 34 of 100, B. 2., after "2. Food Safety and Foreign Material:", delete text up to "3. Container Integrity Defects:" in entirety and insert:
- (a) All corrective actions performed on product due to foreign material and/or processed/unprocessed container mix-ups must be approved by FTR. FTR approval may be accomplished by means of one the two following methods, the methods being subject to change as determined by the contracting officer to be necessary for determining FTR approval:

METHOD 1:

All corrective actions performed on product due to foreign material and/or processed/unprocessed container mix-ups shall be submitted by the contractor to the GQAR for review and acceptability determination. This requirement only applies to contractor facilities that are producing product and/or placing food product into finished component packaging.

If the GQAR determines that the corrective action plan is acceptable, the contractor shall submit a "foreign material notification" or "unprocessed container notification" to FTR, prior to offering the lot for Government inspection. The notification shall include the corrective action plan, the GQAR's recommendation pertaining to the plan, and supporting documentation. FTR shall issue written authorization for offer of the lot for Government inspection.

If the GQAR determines that the correction actions are not acceptable and GQAR and the contractor cannot agree to an alternate plan for remediation, the contractor shall submit a corrective action/remediation plan and supporting documentation to FTR for resolution.

Standard rework procedures (SRP) for specific foreign material situations may be addressed under the contractor's documented QSP, Section XII - Corrective and Preventive Action Program. (see E-4-G.B.6.). SRP's shall only be submitted to DLA for foreign material inherent to a specific food product or ingredient. Screws, plastic pieces, bandages, metal fragments, glass, etc., are not inherent to ingredients used in food products and a SRP for these types of foreign material shall not be submitted to DLA for consideration and approval. SRPs submitted to DLA for review and approval shall have a title beginning with "Standard Rework Procedure for...". SRPs may be referenced, as applicable, in the corrective action plan that he contractor provides for a specific instance (along with any relevant specific details).

All preventive and corrective actions documented by, proposed by, and conducted by the contractor shall conform to the regulations promulgated by the applicable regulatory agency (FDA, USDA-FSIS, USDC). When a contractor is required by regulation to notify a regulatory agency regarding foreign material and/or processed/unprocessed container mix ups, it shall be the responsibility of the contractor to present to the GQAR and DLA verification of conformance to the applicable agency's regulations.

In all cases, it is recommended that the GQAR be notified as soon as possible if and when incidents involving the

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Part 12 Clauses (CONTINUED)

finding by the contractor of foreign material in product and/or product ingredients.

METHOD 2:

The contractor shall submit a corrective action plan and supporting documentation to FTR for resolution.

Standard rework procedures (SRP) for specific foreign material situations may be addressed under the contractor's documented QSP, Section XII - Corrective and Preventive Action Program. (see E-4-G.B.6.). SRPs may be referenced, as applicable, in the corrective action plan that the contractor provides for a specific instance (along with any relevant specific details).

All preventive and corrective actions documented by, proposed by, and conducted by the contractor shall conform to the regulations promulgated by the applicable regulatory agency (FDA, USDA-FSIS, USDC). When a contractor is required by regulation to notify a regulatory agency regarding foreign material and/or processed/unprocessed container mix ups, it shall be the responsibility of the contractor to present to the GQAR and to DLA verification of conformance to the applicable agency's regulations.

In all cases, it is recommended that the GQAR be notified as soon as possible if and when incidents involving the finding by the contractor of foreign material in product and/or product ingredients.

(b) The GQAR shall be notified, and documentation provided, when any finished product intended (or initially intended) to be offered to the Government has been produced using a bulk product or ingredient product lot(s) (or portion thereof) that has, at any time, been identified as containing or having contained foreign material. This requirement only applies to contractor facilities that are producing product and/or placing food product into finished component packaging. The documentation shall identify the foreign material and all corrective actions taken to render the bulk/ingredient product serviceable, including, but not limited to segregation and removal of portions of the bulk/ingredient product. The GQAR shall determine if the corrective actions taken render the bulk/ingredient product serviceable, the contractor shall submit a notification, to include the corrective action plan and supporting documentation, to FTR prior to offering any related finished product lots for Government inspection.

When the GQAR determines that the actions taken do not render the bulk/ingredient product to be serviceable and an alternate plan for remediation cannot be agreed upon by the GQAR and the contractor, the contractor shall submit a corrective action plan and supporting documentation to FTR for resolution.

Standard rework procedures (SRP) for specific foreign material situations may be addressed under the contractor's documented QSP Section XII - Corrective and Preventive Action Program (see E-4-G, B, 6). SRP's shall only be submitted to DLA for foreign material inherent to a specific food product or ingredient. Screws, washers, plastic pieces, bandages, metal fragments, glass, etc., are not inherent to ingredients used in food products and a SRP for these types of foreign material shall not be submitted to DLA for consideration and approval. SRPs submitted to DLA for review and approval shall have a title beginning with "Standard Rework Procedure for...". These SRPs may be referenced, as applicable, in the corrective action plan that the contractor provides for a specific instance (along with any relevant specific details).

All preventive and corrective actions documented by, proposed by, and conducted by the contractor shall conform to the regulations promulgated by the applicable regulatory agency (FDA, USDA-FSIS, USDC). When a contractor is required by regulation to notify a regulatory agency regarding foreign material and/or processed/unprocessed container mix ups, it shall be the responsibility of the contractor to present to the GQAR and to DLA verification of conformance to the applicable agency's regulations.

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Part 12 Clauses (CONTINUED)						
In all cases, it is recommended that the GQAR be notified as soon as possible if and when incidents involving the finding by the contractor of foreign material in product and/or product ingredients.						

(c) Thermal process deviations or deviations from the preparation, formulation or critical factors cited in the approved process schedule must be accompanied by a detailed letter from the plant's Processing Authority. The involved subcode(s), the deviation, and the disposition of the product shall be clearly identified when the complete lot is presented for Government end item verification inspection. If the producer fails to provide enough information/data in the case of a deviation, the GQAR shall contact FTRC for approval to proceed with the Government end item verification inspection.

verification inspection. (d) Retesting/reinspection/rework of product that tested positive for food borne pathogens is not authorized. (e) These requirements are in addition to applicable Code of Federal Regulations or other regulatory requirements (USDA-FSIS, FDA). **NOTE:** Deviations (that occur during or prior to the production of a product) from specific preparation/ formulation/ ingredient requirements cited in the specifications shall be submitted as a request for product deviation through the applicable contracting officer for the coordination with and the approval of the Specification Preparing Activity (Natick). All other terms and conditions remain the same