## Sanitation SOP Development

This form has been designed to provide a guide in assessing sanitation needs and procedures for equipment.

SOP Document Number Issue Date Revision/Review Date Equipment Description	Each SOP should have a unique identifier for the site's document management program. It also should have an initial Issue Date and Revision/Review Date.
Intended Use	Choose best choice(s): In-process Ingredient   RTE Non-RTE
Personnel Responsible to Clean Equipment	Depending on the situation, it may be prudent to list responsible parties (i.e. sanitation crew, operations, contractor, etc.).
Frequency of Cleaning Recommendation	
Sanitation Performance Standard to be Achieved <sup>(1)</sup>	Choose best choice(s): Allergen CleanQuality CleanMicro Clean
Cleaning Chemicals Needed	
e.g. concentrations; water temperature; rinse water pressure if applicable (non- CIP)	
Equipment and Tools Needed	
e.g. vacuum (central/portable), brushes, buckets, scrapers, etc	
Personal Protective Equipment Needed ( reference LOTO Procedure)	Choose Best Choice(s):GlovesEye ProtectionFace MaskRespiratorOther:
Pre-Cleaning (gross soil	Step Procedure
removal) / Scrap Procedure	
removal) / Scrap Procedure	1
	2
	3
	4 5
	6

Disassembly	Step	Procedure
	1	
	2	
	3	
	4	
	5	
	6	
Cleaning Steps	Step	Procedure
	1	
Include chemical clean,	2	
mechanical scrubbing and	3	
rinse procedures; time/temp	4	
and flow rates for CIP.	5	
	6	
	7	
	8	
	9	
	10	
	11	
	12	
	13	
Inspection / Verification <sup>(2)</sup>	Step	Procedure
Procedure	1	
	2	
e.g. pre-op inspection; ATP	3	
swab; protein swab; chemical	4	
concentrations; chart review if	5	
applicable; visibly clean using	6	
powerful light; allergen swab;	7	
etc.	8	
	9	
Actions if verification fails (i.e.	10	
reclean equipment).	11	
Assembly	Step	Procedure
	1	
	2	
	3	
	4	
	5	
Final Sanitation Step	Step	Procedure
	1	
	2	
	3	
	4	
	5	

Validation Procedures <sup>(3)</sup>	Step	Procedure
	1	
e.g. verification results over	2	
time; ATP swab; protein swab;	3	
ELISA allergen test; chemical	4	
concentrations; chart review if	5	
applicable, etc.	6	
Actions if validation fails (i.e. review cleaning procedure).	7	
	8	
	9	

Allergen Clean - Reduction or elimination of residual allergen-containing products.
Quality Clean - Visibly clean.
Micro Clean - Reduction or elimination of target organisms (can be spoilage or pathogenic

organisms depending on risk).

(2) Verification - those activities conducted to prove or substantiate that the sanitation procedures are being properly implemented to achieve the desired objective.

(3) Validation - A review of verification data over a period of time to determine if the procedure consistently produces expected results (e.g. cleaned right the first time). Trending data that shows repeated failures means a review of the procedures and training needs to be conducted.

## **Other Considerations:**

- Note where cleaning is to occur. In process room or designated cleaning areas? If designated cleaning areas, what steps are necessary to prevent recontamination while moving back to processing room?
- Evaluate the need for additional sanitizing steps to ensure all surfaces are sanitized. Sanitizing steps may be a part of pre-assembly and assembly as well as after final rinse.
- There may be a need to perform a post-assembly verification of sanitation to ensure the equipment is not re-soiled during assembly.