Process evaluation

Lethality

- Containers do not heat instantaneously
- All temp. have a lethal effect above 93°C
- Contributes to the destruction of m.o.
- Mechanism to determine the relative effect of a changing temperature is necessary
- z value helps to calculate the lethal effect of various temperature

Lethal Rate (L)

- The relative effect of temp. on microbial destruction with respect to a certain reference temp. (Tref)
- Equivalent time in min at reference temp / time in min at temp. (T)
- L = 10 (T-Tref)/z

Lethal Rate (L)

Temp (oC)	Lethal rate	Time (F _T) req.
	(min at 121.1oC/ min at T)	for 12 D
101.1	0.01	4 h
111.1	0.1	24 min
121.1	1	2.4 min
131.1	10	15 s
141.1	100	1.5 s

Change in processing temp by one z value requires 10-fold change in processing temp



Figure 2 Thermal death time curve passing through 1 min at 121.1 °C.

Sterilization value

- Lethal effect as a function of time (t) during the thermal process
- $\blacksquare F_{\text{Tref}} = {}_0 \int^1 \mathbf{10} \ (\text{T-Tref})/z \ dt$
- $\blacksquare \mathsf{F}_{\mathsf{Tref}} = \Sigma \mathsf{L} \Delta \mathsf{t}$
- When T is the cold point temp. and when ref. temp and z value are 121.1°C and 10°C, then, the sterilization value is known as Fo value
- Fo value of 3 min is acceptable.

Commercial Sterility

Application of heat that render the food free from viable forms of m.o of public health significance as well as any m.o of non PHS capable of reproducing under nonrefrigerated conditions of storage and distribution.

Process evaluation

- Sterilizing effect : Starts at temp. above their maximum growth temp.
- Lethal effect on bacterial spores commence at 93°C
- Process time Time theoretically required to destroy any specific spore in food can at any given temp is calculated from
 - Lethal rate of the temperature
 - TDT curve
 - Heating and Cooling curves

Thermal process evaluation

Determination of thermal process, F_o involves
Measurement of temp. at SHP
Integration of lethal effect at this point

Lethality

- Lethal rate Rate of destruction of an organism per min at any given lethal temp. is the reciprocal of the time in min required to completely destroy it at that temp. LR = Antilog [(T-121.1)/z]
 - If z value is 10°C, the LR for 111.1°C is 0.1



Integration of lethal effects

- C. botulinum spores requires F_o of 2.52 min
 - Heat the food at 121.1°C hold for 2.52 min instantaneously, and cool below lethal temp.
- Integrated lethal value equivalent to 2.52 min sufficient
- F_o at 121.1°C is unity
 - So, for every temp. change of 10°C, there is 10-fold change in death rate of organisms
 - Work out F_o equivalent at that time-temp combinations
 - If processing at 111.1°C, needs 10 times more heating to attain sterilization effect
 - If processing at 131.1°C, needs 1/10th of time



- Methods of estimating process lethality
 - Classical calculation method
 - General method
 - Formula method
 - Integrated lethality method
 - Nomogram method

Classical calculation method

- Simplest method
- Record SHP temp/ min
- Obtain LR rate for temp. from LR table or calculate LR
- Integrate lethal values of heat for all time-temp combinations at this point
- Cumulative F_o value Add LR of temp after steam cut off also

General method (Bigelow, 1920)

- Graphical method each point of H & C curve represent lethal value for the organisms
- Construct TDT, H&C curves and lethality curve with LR against time (min)
- Area under lethality curve gives the total lethality of the process i.e summation of all lethal rates obtained during CUT, retorting and cooling.
- Measure area by counting squares or using planimeter

General method (Graphical method)

- If area under lethality curve is unity (1) process adequate – complete destruction achieved
- If total lethality < 1 under processing</p>
- If total lethality > 1 over processing
- Process value (Fo)
 - = Area under LR curve x lethality/square
 - = 117 x 0.1 = 11.7 min
- Cannot predict F_o of shorter or longer process, change in retort temp. or can size

Mathematical method (Ball and Olson, 1928)

- Applicable to convection and conduction packs of any can size and retort temp
- Construct HP curve by plotting values representing differences between retort temp. and product temp. on a log scale and time on linear scale
- Zero time taken when the steam is turned on
- After an initial lag, the graph becomes a straight line

Mathematical method

Slope is fh

- CUT sterilizing effect is 42% of CUT
- Corrected zero time is 52% of CUT
- Extend straight line of graph to meet 'y' axis
- The corresponding temp is theoretical initial temp (I)
- Ratio of jl/l is referred to as 'j'
 - jl = diff. between retort temp. and theoretical initial temp
 - I = diff. between retort temp and actual initial temp
 - j = lag factor or time before which there is no increase in temperature at SHP

Mathematical method

The basic equation is

B process time (min) = fh log (jl-g)

- g = diff. in oF bet. retort temp and max. temp at SHP
- From above, g is calculated
- Value of g is related to the ratio of fh/U
 - $U = F_o F_i$
 - Fi = Time in min at the retort temp equivalent to F_o process at 250oF
 - Fi = log -1 (250-t)/z (t process temp; z-18oF)
- From above, Fi is calculated
- From U and Fi, Fo is calculated

Integrated lethality method

- In convection pack, heating at SHP (Fc) is equal to the effective mean through out the container (Fs)
 - Fc = Fs = D (log a log b)
 - **D** = **DRT** at reference temp
 - a = initial no of spores
 - b = no of survivors
- For conduction pack, integration of lethal values of heat received at SHP is more accurate
 - Determine survival ratios for small volume
 - Integrate over the whole volume

Nomogram method (Olson and Stevens, 1939)

- American Can Co Quick method
- Not accurate
- Applicable to straight line and broken curve lines
- Knowledge on Ball method necessary
- Calculate process value (Fo)

Predict Fo values for similar products under changed conditions ie. can size, retort temp or filling temp.

Verification of the Process

Microbiological methods

- Most reliable
- Two methods
 - Inoculated pack method Spores are directly added to the food
 - Spore bulb method Spores suspended in a buffer and contained in a small bulb or capillary tube are placed in the food in the can

Inoculated pack method

- Place heat resistance spores (gas former) in SHP
- Subject to normal heat treatment
- Presence of blown can
- Estimate survival by culture or incubation tests

Spore bulb method

- Similar to capillary bulb method
- Capillaries containing spores implanted in solid pieces of food at SHP
- In convection packs, placed using special device
- Compare no. of survivors with the initial no. of spores
- Evaluate the lethality of the process