Table 2. Parameters of length distributions obtained from batch- and chemostat-cultured cells, prepared by agar filtration

Culture	τ (min)	\hat{L} (μ m)	CV ^a (%)	L_0^b (μ m)	Cells showing con- striction			Cell width (µm)	
					%	L_c^c $(\mu \mathbf{m})$	CV (%)	2R (μm)	SD ^d (µm)
B/r A				-					
Batch ^e	160	$1.4 (1,025)^{f}$	23	1.0	14	1.9	8	0.65	$0.04 (144)^f$
Batch ^o	126	1.6 (632)	24	1.1	14	2.2	8	0.64	0.03 (69)
Chemostat	175	1.5 (974)	23	1.1	9	2.1	8	0.56	0.04 (112)
B/r K		, ,							
Batche. g	180	2.2 (1,140)	24	1.6	6	3.1	11	0.46	0.04 (119)
Chemostat	175	2.4 (955)	25	1.6	8	3.2	15	0.52	0.04 (127)

^a CV, Coefficient of variation.

^b The length of newborn cells (L_0) was estimated from the distributions using the formula (11): $L_0 = \frac{1}{2}$ $(L_{\min} + \frac{1}{2} L_{\max})$, where L_{\min} and L_{\max} are, respectively, the minimal and maximal cell lengths observed.

^c L_c , the average length of cells showing constriction, is usually somewhat smaller than $2 \times L_0$.

^d SD, Standard deviation.

^e See Fig. 1 for length distributions.

^f The number in parentheses indicates the number of cells measured.

⁹ Cultures used for analysis of nuclear separation in critical-point-dried cells (cf. Table 4).