## PRODUCT SPECIFICATION



## CMA 003 HybER–Zero Hybridoma Enhancing Reagent, serum free

Culture Medium Additive Lyophilized

Article No.	71800		
Product Name	CMA 003 HybER-Zero Hybridoma Enhancing Reagent, serum free		
Presentation	Appearance: (	Clear light yellow or orange	solution.
	Content: a	2.5 mL, lyophilized	
Storage	Lyophilized: a	at room temperature	
		at -18°C or colder for long-	term storage
•••••		at 2-8°C when in use	
Expiry		Exp.date	
		Exp.date at -18°C or colder	
		3 months after reconstitut	ION at 2-8°C
Sterility		Non-sterile	
		yophilized and sealed unde	r aseptic conditions
	•	Negative in PCR screening Negative in LAL-test	
Decemintion	· · · · · · · · · · · · · · · · · · ·	· · • · · · · · · · · · · · · · · · · ·	mas immediately after fusion and
Description	HybER-Zero stimulates growth of mouse hybridomas immediately after fusion and during cloning procedures (1,2). HybER-Zero has been especially formulated for		
	serumfree hybridoma o		
Protocol of use	To prepare HybER-Zero for laboratory use:		
	1) Reconstitute with 2.5 mL water for injection (WFI) or culture medium directly into the		
	vial with lyophilized HybER-Zero.		
	2) Solubilise all material by pipetting gently up and down.		
	3) Filtrate the reconstituted HybER-Zero through a 0.22 $\mu m$ sterile filter.		
	HybER-Zero is now ready for use and suitable amounts can be added as needed to the		
	••••••••••••••••••••••••	um during fusions and cloni	•• <del>•</del> •••••••••••••••••••••••••••••••••
Dilution Guide	We recommend using HybER-Zero at a dilution of 0.5% $(v/v)$ in growth medium immediately after fusion, at the first medium change after fusion, and during		
	immediately after fueir	n at the first medium char	are after fusion and during
	•	-	<b>o</b>
	subsequent cloning ste	-	nge after fusion, and during ilution, one vial of HybER-Zero (2.5 mL)
Application	subsequent cloning ste	eps. At our recommended d	<b>o</b>
Application	subsequent cloning ste is sufficient to supply 5	eps. At our recommended d 500 mL of growth medium.	ilution, one vial of HybER-Zero (2.5 mL)
Application	subsequent cloning ste is sufficient to supply 5 Method	eps. At our recommended d 500 mL of growth medium. Usability	ilution, one vial of HybER-Zero (2.5 mL) Dilution guide
Application	subsequent cloning ste is sufficient to supply 5 Method Fusion	eps. At our recommended d 500 mL of growth medium. Usability yes	ilution, one vial of HybER-Zero (2.5 mL) Dilution guide 0,5% (v/v)
Application	subsequent cloning ste is sufficient to supply 5 Method Fusion Cloning	eps. At our recommended d 500 mL of growth medium. Usability yes yes	ilution, one vial of HybER-Zero (2.5 mL) Dilution guide 0,5% (v/v) 0.5% (v/v)
Application References	subsequent cloning ste is sufficient to supply 5 Method Fusion Cloning Production 1) Trier NH, Mortensen A, Sc	eps. At our recommended d 500 mL of growth medium. Usability yes yes nd. hiolborg A, Friis T. Production and	ilution, one vial of HybER-Zero (2.5 mL) Dilution guide 0,5% (v/v) 0.5% (v/v)
	subsequent cloning ste is sufficient to supply 5 Method Fusion Cloning Production 1) Trier NH, Mortensen A, Sc Methods Mol Biol. 2015;1348	eps. At our recommended d 500 mL of growth medium. Usability yes yes nd. hiolborg A, Friis T. Production and 109-26. J, Houen G. New tools for studying	ilution, one vial of HybER-Zero (2.5 mL) Dilution guide 0,5% (v/v) 0.5% (v/v) nd.

## Conditions

For research use only. Not for use in diagnostic procedures. Not for therapeutic use or applications.

The information and product are offered without guarantee as the ultimate conditions of use are beyond our control. The foregoing is in lieu of all warranties, expressed or implied, including implied warranties of merchantability and fitness for a particular purpose. In no event shall Statens Serum Institut be responsible for loss of profits or indirect consequential losses resulting from use of its products. The animals from which this product was derived have not been exposed to or inoculated with any livestock or poultry disease agents exotic to the United States or Western Europe, and did not originate from facilities where work with exotic disease agents affecting livestock or avian species is carried out.

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