Technical Information

Stepan

Stepan Company
Northfield, Illinois 60093
Telephone 847 446 7500



MAMMALIAN TOXICOLOGY OF DIAMIDOAMINE QUATERNARIES

Applicable to these current Stepan products:

ACCOSOFT® 440-75	ACCOSOFT® 460 HC	ACCOSOFT® 501
ACCOSOFT® 501 DEG	ACCOSOFT® 540	ACCOSOFT® 550-75
ACCOSOFT® 550-90 HF	ACCOSOFT® 550-90 HHV	ACCOSOFT® 550-PG
ACCOSOFT® 550 HFC	ACCOSOFT® 550L-90	ACCOSOFT® 580
ACCOSOFT® 620-75	ACCOSOFT® 620-90	ACCOSOFT® 750
ACCOSOFT® 780	ACCOSOFT® 780 PG	

Applicable to these inactive Stepan products:

ACCOSOFT® 502	ACCOSOFT® 540 HC	ACCOSOFT® 570
ACCOSOFT® 570 HC		

Toxicological Information:

Test/Conditions	Results/Classification	References
Mammalian Toxicology:		
Acute Oral Toxicity	LD ₅₀ > 5g/kg	Stepan Study
(rat) (gavage) (14 day)	(practically non-toxic orally)	No. 78-032B
	, ,	(Industry Consortium Data)
Acute Percutaneous absorption	LD ₅₀ > 2g/kg	Stepan Study
(rabbit) (14 day)	(slightly toxic dermally)	No. 81-015A
	No signs of systemic toxicity	(Industry Consortium Data)
Primary Eye Irritation	$MMS^{1} = 1/110$	Stepan Study
(rabbit) (24 hr)	(minimal eye irritation	No. 78-032C
n=6	@ 5% dispersion)	(Industry Consortium Data)
Primary Skin Irritation	$PII^{2} = 0.8/8$	Stepan Study
(rabbit) (24 hr exposure)	(slightly irritating to skin @	No. 78-032C
n=6	5% dispersion)	(Industry Consortium Data)
Human Patch Test	Minimal skin irritation at	Industry Consortium Data
(24 hr. exposure)	concentrations of up to	
(over a 6 day period)	20% w/v	
Human Patch Test	No skin sensitization	Industry Consortium Data
(24 hr. contact) (9 exposures	observed 25% w/v aqueous	

over 21 days) (n=205)	solution	
Subchronic Percutaneous Toxicity Study (rabbit) (4 wks)	No treatment related chemical changes observed. The systemic no-observed effect level (NOEL) was 300mg/kg	Industry Consortium Data
Genotoxicity Studies		Industry Consortium Data
i.) Ames	Not a mutagen	
ii.) Mouse Lymphoma Assey	Negative	

MMS¹=Maximum Group Mean Score PII² = Primary Skin Irritation Index

References:

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