

RINSKORTM ACTIVE MAMMALIAN TOXICOLOGY

RinskorTM active has a very favorable mammalian toxicity profile.

Rinskor and its formulated products exhibit low acute toxicity with mild eye irritation being the most significant acute findings with the technical material as well as the formulated product. No significant adverse effects were reported in the sub-chronic, chronic, reproduction or developmental toxicity studies, nor were there any treatment-related mutagenic, carcinogenic or neurotoxic effects noted. Mammalian tox profile is described below.



Selected Mammalian Toxicological Endpoints

Study	Results
Acute oral, rat	LD ₅₀ >5000 mg/kg
Acute dermal	LD ₅₀ >5000 mg/kg
Acute inhalation	LC ₅₀ >5.23mg/L
Eye irritation	None
Skin irritation	None
Skin sensitization (LLNA)	Weak dermal sensitization (EC ₃ =19.1%)
Chronic Toxicity/Carcinogenicity: Rats	No evidence of long term toxicity or carcinogenicity up to the highest dose tested, 300 mg/kg bw/day. Chronic NOAEL= 300 mg/kg bw/day
Carcinogenicity: Mice	No evidence of carcinogenic potential. NOEL/NOAEL = 1000 mg/kg/day (M)/800 mg/kg/day(F)
Developmental Toxicity: Rat	No evidence of developmental toxicity up to the limit dose. Maternal NOAEL= 1000 mg/kg/day Litter NOAEL = 1000 mg/kg/day
Developmental Toxicity: Rabbit	No evidence of developmental toxicity up to the limit dose. Maternal NOAEL= 1000 mg/kg/day Litter NOAEL = 1000 mg/kg/day
2-Generation Reproduction: Rat	No evidence of reproductive toxicity up to the highest dose tested, 300 mg/kg/day. Parental NOAEL= 300 mg/kg/day Pup NOAEL = 300 mg/kg/day



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RinskorTM active registrations are pending in several countries around the world. See the country-specific information on this site. Always read and follow label instructions.