1 物質番号

<u>' 1775 A 7</u>	
通し番号	C-1001
整理番号	1
MITI番号	
	51-03-6
物質名	ピペロニルブトキシド
英名	Piperonyl butoxide

2 発がん性分類

2	<u> 心性が親</u>		
機関名	分類 結果	評価年	評価書引用文献
IARC	3	1987	 National Technical Information service (1968) Evaluation of carcinogenic, Teratogenic and mutagenic Activities of selected pesticides and Industrial Chemicals, Vol. 1, Carcinogenic study (PB−223159) National cancer Institute (1979) Bioassay of Piperonyl Butoxide for possible carcinogenicity (Tech. Rep. ser. NO. 120; DHEW publ No.(NIH)79−1375) Sarles, M.P. & vandegrift, W.B. (1952) chronic oral toxicity and related studies on animals with the insecticide and pyrethrum synergist, piperonyl butoxide.Am. J. trop. Med. Hyg., 1, 862−883 Epstein, SS. Joshi, S., Andrea, J, Clapp, P., Falk, H. & Mantel, N. (1967) synergistic toxicity and carcinogenicity of 'Freons' and piperonyl butoxide. Nature, 214, 526−528
EPA	×	ı	-
NTP	×	ı	-
ACGIH	×	-	_
産衛学 会	×	-	-
EU	×	_	-

3 発がん性に関する追加文献(動物試験、疫学調査)

追加文献の有無 有

(1)動物試験

<u>(1)動物</u> 	試験概要	訂	式験物質	試験の種類	ガイドライン	GLP適用状況	試験 実施年	試験 実施者	
				chronic test			1997	Tokyo Metropolitan Research Laboratory of Public Health	
	試験条件	動物種	系統	動物数/ 性別/群	投与経路	用量/濃度	単位	投与/暴露期間	
#1		mouse	Grj:CD−1	control: 52(M)/51(F), 0.6%diet:53 (M)/52(F), 1.2%diet: 100(M)/106(F)	feeding	0, 0.6, 1.2	%	52 weeks	
	試験結果概要	発がん影響	related manner in the 1.2% gro	in males, with a	a 5–fold increas sex difference	e induced by piperonyl but e with a doubling of the do was observed in hepatoca n males.	ose, while	e they were found only	
		非発がん影響	hemorrhage of	Kidneys were black-colored in several treated male mice and were dose-dependent. Edema and nemorrhage of ovary were observed in all groups of female mice. Hemorrhages in many organs were found especially in the dead male mice of the 1.2% group.					
		結論		xide is a hepato	_				
	文献名		akahashi , S. Oishi, T. Fujitani, T. Tanaka, M. Yoneyama , "Chronic toxicity studies of piperonyl butoxide ir 1 mice:Induction of hepatocellular carcinoma" Toxicology 124 (1997) 95–103						

		訂	式験物質	試験の種類	ガイドライン	GLP適用状況	試験 実施年	試験 実施者			
	試験概要			chronic toxicity study			1994	Tokyo Metropolitan Research Laboratory of Public Health			
		動物種	系統	動物数/ 性別/群	投与経路	用量/濃度	単位	投与/暴露期間			
#2	試験 条件	mouse	Crj:CD-1	male control: 52, 0.6%diet: 53, 1.2%diet: 100	diet	0, 0.6, 1.2	%	12 months			
	= →F 2 &±	発がん 影響	control groups.	. The incidences	s of hepatocellu	ted groups in a dose-de lar carcinoma were 11.3 nyl butoxide can cause l	and 52.0%	in mice given 0.6 and			
	試験結果概要	非発が ん影響									
		結論	In conclusion, to carcinoma in m	•	ults clearly indic	ate that Piperonyl buto	xide can in	duce hepatocellular			
	文献名	Takahashi O, Oishi S, Fujitani T, Tanaka T, Yoneyama M "Piperonyl butoxide induces hepatocellular carcinoma in male CD-1 mice." Arch Toxicol. 1994;68(7):467-9.									
		ā	式験物質	試験の種類	ガイドライン	GLP適用状況	試験 実施年	試験 実施者			
	試験 概要			chronic toxicity study			1994	Tokyo Metropolitan Research Laboratory of Public Health			
		動物種	系統	動物数/性別/群	投与経路	用量/濃度	単位	投与/暴露期間			
	試験条件	rat	F344	30 animals/dose/ sex, 2.4%: 33 animals/sex	diet	0, 0.6, 1.2, 2.4	%	nearly 2 years			
#3		発がん影響									
	試験結 果概要	非発がん影響	Piperonyl buto Hemorrhages i	Beginning at about 40 weeks, 10 rats in the 1.2% treated male group died due to cecal hemonoperonyl butoxide also caused essential thrombocythemia with a dose-response relationsh Hemorrhages in stomach and cecum, anemia, degenerative lesions of alveoli, and nephrotoxical observed related to exposure.							
		結論	These results i	indicate that pip	peronyl butoxide	is a hepatocarcinogen	to the rat.				
	文献名					M. ″Chronic toxicity stu dam Appl Toxicol. 1994					

	試験概	ā	式験物質	試験の種類	ガイドライン	GLP適用状況	試験実 施年	試験実施者		
	要			Carcinogenicit y Studies			1979			
		動物種	系統	動物数/性別/群	投与経路	用量/濃度	単位	投与/暴露期間		
#4	試験条件	rat	F344	control: 20 animals/sex, 50 animals/dose/ sex	feeding	5000, 10000	ppm	2 years		
	=4 F4 4+	発がん 影響	was associated	l with administra		d increase in the incidence opound to females, the inc treated males.				
	試験結 果概要	非発が ん影響								
		結論	_	_		ex is not considered by its etment of carcinogenic ac		nstitute sufficient		
	文献名	Carcinogenesis bioassay of technical-grade piperonyl butoxide in F344 rats. Cardy RH, Renne RA, Warner JW, Cypher RL J Natl Cancer Inst. 1979 Mar;62(3):569-78								
		試験物質		試験の種類	ガイドライン	GLP適用状況	試験実 施年	試験実施者		
	試験概要			Combined Chronic Toxicity / Carcinogenicit y		yes	1998	Bio-Research Ltd Laboratory		
	= 0 5 0 47	動物種	系統	動物数/性別/群	投与経路	用量/濃度	単位	投与/暴露期間		
#5	試験条 件	rat	Sprague- Dawley	60/sex/dose	feeding	0, 30, 100, 500	mg/kg/ day	104/105weeks		
		発がん影響	There was no i	ncreased incide	nce of neoplasi	a at any site.		1		
	試験結果概要	非発がん影響		nd female rats.		500 mg/kg/day associate d hyperplasia of thyroid fo				
		結論			nce of neoplasi	-				
	文献名		H, Gabriel KL, C Toxicol. 1998 (-	renicity studies of piperon	yl butoxid	de in rats and mice."		

	= 45 45 100	ā	式験物質	試験の種類	ガイドライン	GLP適用状況	試験実 試験実施者				
	試験概 要			Carcinogenicit y Studies		yes	1998	Bushy Run Research Center,			
	- > = > =	動物種	系統	動物数/性別/群	投与経路	用量/濃度	単位	投与/暴露期間			
"0	試験条件	mouse	CD1	60/sex/dose	feeding	0, 30, 100, 300	mg/kg/ day	78weeks			
#6		発がん影響				e was evidence of increas at 100 and 300 mg/kg/da					
	試験結 果概要	非発が ん影響									
		結論	oxygenase grou		In the mouse th	es related to the induction e increased incidence of e					
	文献名	Butler WH, Gabriel KL, Osimitz TG, Preiss FJ. "Oncogenicity studies of piperonyl butoxide in rats and mice." Hum Exp Toxicol. 1998 Jun;17(6):323–30									
	試験概要	詞	式験物質	試験の種類	ガイドライン	GLP適用状況	試験実 施年	試験実施者			
		Technica	al-grade BP	Carcinogenicit y Studies			1985	National Institute of Hyginic Sciences			
	試験条件	rat	F344	46-49 animals/dose/ sex	diet	0, 0.5, 1	%	2 years			
#7		発がん 影響 Various tumours were detected in all groups, including the untreated control group, but no dose-related increase in the incidence of any tumour was found.						oup, but no significant			
	試験結 果概要	非発がん影響	Ileocaecal ulcers were found in animals of both sexes in both experimental groups and the incidence was dose related.								
		結論	F344 rats.		·	conditions piperonyl buto		_			
	文献名	technica				Hayashi Y. "Lack of evide ive induction of ileocaecal					

	=-1: E-0 100	試験物質		試験の種類	ガイドライン	GLP適用状況	試験実 施年	試験実施者	
	試験概 要			combined chronic/carci nogenic study					
	試験条	動物種	系統	動物数/性別/群	投与経路	用量/濃度	単位	投与/暴露期間	
	件	rat and mouse							
#8	試験結果概要	発がん	where a high in carcinomas we thyroid follicula follicular cell tu study reported	n a combined chronic/carcinogenic study in rats, positive carcinogenic effects were reported at doses where a high incidence of ileocecal ulcers were noticed in test mammals. Liver adenomas and carcinomas were reported in Fischer 344 rats only when tested at very high doses. A slight increase in chyroid follicular cell tumors was reported in Sprague-Dawley rats. A slight increase in thyroid follicular cell tumors was reported in Sprague-Dawley rats. A 1979 National Toxicology Program (NTP) study reported negative effects for carcinogenicity in the same strain of rats and in B6C3F1 mice. In CD-1 mice, PBO tested positive for liver tumor effects.(EPA REDより引用)					
		非発が ん影響							
		結論	PBO is classifie	ed as a Group C	C -possible hum	an carcinogen(EPA RED↓	り引用)		
	文献名	EPA REI)						

(2)疫学調査

#1	調査の 種類	調査方法	結果の概要	調査実 施年	調査実施者
"'	文献名				