

[4] 酸化プロピレン

試験法	試験方法	使用生物種・細胞株	試験結果		文献番号	
			代謝活性化系			
			あり	なし		
in vitro	SOS 修復	ネズミチフス菌 TA1535/pSK1002	+	+	1	
	前進突然変異	バクテリオファージ		-	2	
	復帰突然変異	ネズミチフス菌 TA100		(+)	3	
			+	+	4	
			+	+		
			+	+		
				-		
				+	6	
				+		
				+		
				+		
			+			
	ネズミチフス菌 TA1535	ネズミチフス菌 TA1535	-	-	8	
			+	+	4	
			+	+		
			+	+		
				+	6	
				+		
				+		
				+		
				(+)		
			+	-	8	
	ネズミチフス菌 TA1537	ネズミチフス菌 TA1537	-	-	4	
			-	-		
			*	*	9	
	ネズミチフス菌 TA1538		-	-	10	
	ネズミチフス菌 TA98	ネズミチフス菌 TA98	-	-	4	
			-	-		
			-	-		
				-	6	
				+	9	
	ネズミチフス菌 TA97	ネズミチフス菌 TA97		-	8	
				+	9	
	大腸菌 WP2 uvrA	大腸菌 WP2 uvrA	+	+	4	
				-	5	
	大腸菌 WP2	大腸菌 WP2	+	+	4	
			-	+	10	
	大腸菌 B(Arg-)Hs30R			+	11	
	酵母菌			+	13	
	アカパンカビ			+	15	
前進突然変異	肺炎桿菌			+	12	
	酵母菌		+	+	14	
遺伝子交換	酵母菌			+	13	
遺伝子突然変異	チャイニーズハムスター卵巣細胞 , <i>hprt</i> 座位			+	16	
	マウスリンパ腫 L5178Y 細胞, <i>tk</i> 座位			+	17	
染色体異常	チャイニーズハムスター卵巣細胞		+	(+)	18	

		ラット肝細胞	+	19
			+	10
		ヒトリンパ球	+	4
姉妹染色体分体交換	チャイニーズハムスター卵巣細胞	+	18	
	チャイニーズハムスターV79 細胞	+	20	
	ラット肝細胞	+	19	
	ヒトリンパ球	+	21	
DNA一本鎖切断	ラット肝細胞	+	22	
in vivo	伴性劣性致死突然変異	ショウジョウバエ	+	23
	小核誘発	マウス骨髄細胞	+	4 他
			+	
			-	4
	ヒトリンパ球		*	24
	染色体異常	サルリンパ球	-	25
		マウス骨髄細胞	+	26
		ヒトリンパ球	*	24
	優性致死	マウス	-	2
		ラット	-	23
	姉妹染色体分体交換	サルリンパ球	-	25
		マウス骨髄細胞	+	26
	共有結合	マウスのDNA	+	27
		ラットのDNA	+	28
		マウス、ラット及びイヌのDNA	+	29
	精子形態異常	マウス	-	23
	DNAとの共有結合	仔ウシ胸腺DNA	+	30 他
			+	
			+	
評価結果	タンパク質との共有結合	マウス	+	27
		マウス、ラット及びイヌのヘモグロビン	+	29
		ヒトのヘモグロビン	+	31

注：1) + 陽性 ; (+) 弱い陽性 ; - 陰性 ; * 結論が出なかったもの

空欄；試験系がないか、試験されなかつたもの

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