

检测报告 报告编号: ASH20-007421-01 发布日期: 2020-04-22

客户名称: 江西浣星谷科技有限公司

客户地址: 江西省赣州市章贡区橙乡大道 26 号嘉福大厦 1909 室

样品名称: 天然矿物功能剂

样品批号: 20200108 生产日期: 2020/1/02

生产商: 江西浣星谷科技有限公司

样品其他信息: 纳米功能剂

以上样品及信息由客户提供及确认, SGS 不承担证实客户提供信息的准确性、适当性和(或)完整性责任。

样品接收日期: 2020-03-09

检测周期: 2020-03-09 ~ 2020-04-22 检测要求: 根据客户要求进行检测

检测方法: 请参见下一页 检测结果: 请参见下一页

除非另有说明,本检测结果仅与被检测物品有关。仅供客户内部使用,不对社会具有证明作用。未经检验机构书面同意,委托人不得擅自使用检测结果进行不当宣传。







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检测样品描述:

样品编号 SGS 样品 ID 描述

1 ASH20-007421.001 灰色粉末

检测方法:

1.*皮肤刺激试验: 16 CFR 1500.41

检测环境: 普通动物房,实验动物使用许可证号: SYXK(浙)2018-0003,室温 21.5℃~23.0℃; 相 对湿度 52.9%~64.8%。

试验动物: 新西兰白兔,由桐乡市银海牧业专业合作社提供,生产许可证号: SCXK(浙)2018-0002。 动物数/性别: 6 只, 雌雄各半 (雌性动物未孕和未曾产仔)

试验方法:实验动物单笼饲养,试验前在实验动物房环境中至少适应 3d。试验前 24h 去除动物脊柱两侧 被毛,去毛范围左、右各约 2.5cm×2.5cm,选择皮肤健康完整无损的动物进行试验。涂受试物前,在 每只动物的右侧去毛皮肤上,用75%酒精清洁、消毒暴露皮肤,待酒精挥发后,用灭菌刀片或注射针 头在皮区内划一个"井"形的破损伤口。取 0.5g 原样品用生理盐水湿润后涂抹在两层纱布(约 2.5cm ×2.5cm 大小)上并贴敷于两侧去毛区,用一层玻璃纸覆盖,再用无刺激性胶布加以固定。采用封闭性 试验,敷用时间为 24h。

观察时间: 观察移除受试物后 24h 和 72h 时贴敷部位皮肤反应,并进行皮肤反应评分。

2.*眼刺激试验: 16 CFR 1500.41

检测环境: 普通动物房,实验动物使用许可证号: SYXK(浙) 2018-0003, 室温 21.5℃~23.0℃; 相 对湿度 52.9%~64.8%。

试验动物: 新西兰白兔,由桐乡市银海牧业专业合作社提供,生产许可证号: SCXK(浙)2018-0002。 动物数/性别:6只,雌雄各半(雌性动物未孕和未曾产仔)

试验方法: 试验前 24h 对试验动物的两只眼睛进行检查(包括使用 2%荧光素钠溶液检查),确保动物 眼睛可以用于试验。取 0.1mL 原样品投入一侧眼睛结膜囊中,眼闭合 1s。另一侧眼睛不处理作为自身 对照。

观察时间: 入眼后 24 h, 48 h, 72 h。

3.*急性经口毒性试验: GB 15193.3-2014

检测环境: 屏障环境动物房,使用许可证号: SYXK(浙) 2018-0003,温度 21.6℃ \sim 24.1 \textdegree 、相对湿 度 49.2%~55.1%。

试验动物: ICR 小鼠,由浙江省医学科学院实验动物中心提供,生产许可证号: SCXK(浙) 2019-

饲料来源:由江苏省协同医药生物工程有限责任公司生产,生产许可证号为苏饲证(2019)01008,保 质期为6个月,生产日期为2019.12.02,合格证号为120191202008。

动物数/性别: 20 只, 雌雄各半(雌性动物未孕和未曾产仔)

样品制备: 称取 10.0613g 样品置于烧杯中取少量纯水将样品混匀后转入 20mL 容量瓶内,以少量纯水 多次冲洗烧杯一并转入容量瓶中,加入纯水定容至刻度线,充分摇匀后转入试剂瓶标识备用,样品现配 现用。

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试验方法: 采用最大限量法,灌胃剂量为 10061.3mg/kg; 试验前禁食 4 小时。试验开始后,对动物用 经口灌胃法一次染毒,染毒后继续禁食1小时,观察并记录染毒过程和观察期内动物的中毒和死亡情 况,观察周期14天,观察期结束后,处死存活动物并进行大体解剖。

检测结果:

1.原发性刺激积分为0。

附主 4 由胜制游后应任用

动物	动物	は重	红斑				水肿				皮肤	原发		
奶粉 编号	性别	1		破损皮肤		分项	完整皮肤		破损皮肤		分项	反应	刺激	
姍与	土力	(kg)	24h	72h	24h	72h	积分	24h	72h	24h	72h	积分	总分	积分
1	3	2.76	0	0	0	0		0	0	0	0			
2	3	2.42	0	0	0	0		0	0	0	0			
3	2	2.49	0	0	0	0		0	0	0	0			
4	8	2.66	0	0	0	0	0	0	0	0	0	0	0	0
5	2	2.57	0	0	0	0		0	0	0	0			
6	2	2.49	0	0	0	0		0	0	0	0			
皮肤反应评分均值			0	0	0	0		0	0	0	0			

2.在各观察时间(24 h, 48 h 或 72 h),动物结膜、角膜和虹膜的积分均为 0。

附書の眼制激反応は思

附表 2 眼刺激反应结果											
动物			眼刺激性反应积分								
編号	性别	部位	24	4h	48	3h	72h				
細与			样品	对照	样品	对照	样品	对照			
		虹膜	0	0	0	0	0	0			
1	7	角膜	0	0	0	0	0	0			
1	8	结膜充血	0	0	0	0	0	0			
		结膜水肿	0	0	0	0	0	0			
	8	虹膜	0	0	0	0	0	0			
2		角膜	0	0	0	0	0	0			
		结膜充血	0	0	0	0	0	0			
		结膜水肿	0	0	0	0	0	0			
	9	虹膜	0	0	0	0	0	0			
3		角膜	0	0	0	0	0	0			
3		结膜充血	0	0	0	0	0	0			
		结膜水肿	0	0	0	0	0	0			
		虹膜	0	0	0	0	0	0			
4	3	角膜	0	0	0	0	0	0			
		结膜充血	0	0	0	0	0	0			

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=h#m		部位	眼刺激性反应积分							
动物 编号	性别		24	4h	48	3h	72h			
細写			样品	对照	样品	对照	样品	对照		
		结膜水肿	0	0	0	0	0	0		
	9	虹膜	0	0	0	0	0	0		
5		角膜	0	0	0	0	0	0		
9		结膜充血	0	0	0	0	0	0		
		结膜水肿	0	0	0	0	0	0		
)	虹膜	0	0	0	0	0	0		
6		角膜	0	0	0	0	0	0		
0	4	结膜充血	0	0	0	0	0	0		
		结膜水肿	0	0	0	0	0	0		

3.实验动物在染毒 14 天内未见任何中毒症状和中毒死亡;雌雄动物的平均体重未见异常。实验观察结束,对受试动物进行大体解剖检查也未见异常变化。LD50>10061.3mg/kg。

附表 3 急性经口毒性试验结果

사는 단리	动物数		死亡数	死亡率			
性别	(只)	0天	7天	14 天	14 天增重	(只)	(%)
雄性	10	20.4±0.92	24.9±1.08	29.5±1.00	9.1±0.63	0	0
雌性	10	19.9±1.21	23.4±1.33	27.4±1.31	7.5±0.43	0	0

结论:

- 1.根据皮肤刺激强度分级,该样品对兔皮肤无刺激性。
- 2.按眼刺激反应分级,该样品在不冲洗条件下属无刺激性。
- 3. 按照《GB 15193.3-2014 食品安全国家标准 急性经口毒性试验》标准,该样品对 ICR 小鼠的急性经口 $LD_{50} > 5000 mg/kg$ 。根据急性毒性分级,属于实际无毒级。

备注:*检测项目由宁波海关技术中心完成。

*** 结束***

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