

论文

95% 灭菌丹原药对大鼠经口毒性试验研究

柳云恩, 张玉彪, 施琳, 史琳, 张继存, 侯明晓

沈阳军区总医院全军重症战创伤中心实验室及辽宁省重症创伤和器官保护重点实验室, 辽宁 沈阳 110016

摘要:

目的 确定95%灭菌丹原药对SD大鼠的毒性作用,并探寻其可能的毒性靶器官。方法 将80只SD大鼠随机分成4组,各组灭菌丹剂量为0、4 000、8 000及16 000 mg/kg饲料。试验结束后检测动物血常规及生化指标,对解剖动物脏器进行组织病理学评价。结果 试验期间,中、高剂量灭菌丹组大鼠体重均明显降低( $P<0.05$ )。与对照组比较,中、高剂量灭菌丹组大鼠血液中性粒细胞百分率升高,差异均有统计学意义( $P<0.05$ );雄鼠中、高剂量组脑、心脏、脾脏、肝脏、肾脏及肾上腺的绝对重量均高于对照组( $P<0.05$ );中、高剂量灭菌丹组大鼠前胃黏膜下水肿及炎性细胞浸润的发生率均明显高于对照组( $P<0.05$ )。结论 在本试验条件下,前胃是95%灭菌丹潜在的毒性靶器官,该农药的最大无作用剂量为4 000 mg/kg饲料。

关键词: 灭菌丹 靶器官 前胃 黏膜下水肿 暴露风险

Toxicity of 95% captan in rats: an oral toxicity test

LIU Yun-en, ZHANG Yu-biao, SHI Lin, et al

Laboratory of Severe and War-Related Trauma Center of People's Liberation Army, General Hospital of Shenyang Military Region, Shenyang, Liaoning Province 110840, China

Abstract:

Objective To observe the toxic effects of 95% captan and to determine its possible target organ in Sprague Dawley(SD)rats.Methods Eighty SD rats were randomly divided into four groups and administered feeds containing 0(control),4 000,8 000,and 16 000 ppm of captan for 13 weeks.At the end of the study,the blood samples of the rats were collected for hematology and serum biochemistry analyses and tissues of organs were routinely prepared and stained with hematoxylin and eosin for histopathological examination.Results During the experiment,the body weight of the rats in midium and high dose group was significantly lower than that of the control group( $P<0.05$ ),and the percentage of neutrophil was increased with statistical significance compared to those of the control group;the absolute organ weights of brain,heart,spleen,liver,kindey and adrenal glands in high dose group were significantly increased compared to those of the control group( $P<0.05$  for all).The incidence of the edema of forestomach submucosa and inflammatory cell infiltration were significantly higher than those of the control group( $P<0.05$ ).Conclusion Based on the study,forestomach is potential toxic target organ of 95% captan.The maximum no-effect dose of captan is estimated to be 4 000 ppm.

Keywords: captan target organ forestomach submucosa edema exposure risk

收稿日期 2013-09-24 修回日期 网络版发布日期 2013-11-18

DOI: 10.11847/zgggws2013-29-12-27

基金项目:

国家“十一五”重大新药创制资助(2008ZX09305-001)

通讯作者: 侯明晓

作者简介:

参考文献:

[1] Cohen SM, Gordon EB, Singh P, et al.Carcinogenic mode of action of folpet in mice and evaluation of its relevance to humans[J].Crit Rev Toxicol, 2010, 40(6):531-545.

扩展功能

本文信息

- Supporting info
- PDF(913KB)
- [HTML全文]
- 参考文献

服务与反馈

- 把本文推荐给朋友
- 加入我的书架
- 加入引用管理器
- 引用本文
- Email Alert
- 文章反馈
- 浏览反馈信息

本文关键词相关文章

- 灭菌丹
- 靶器官
- 前胃
- 黏膜下水肿
- 暴露风险

本文作者相关文章

- 柳云恩
- 张玉彪
- 施琳
- 史琳
- 张继存
- 侯明晓

PubMed

- Article by LIU Yun-en
- Article by ZHANG Yu-biao
- Article by SHI Lin
- Article by et al
- Article by
- Article by

[2] Gordon E, Cohen SM, Singh P. Folpet-induced short term cytotoxic and proliferative changes in the mouse duodenum[J]. Toxicol Mech Methods, 2012, 22(1): 54-59.

[3] Mireille CR, Beatrice A, Joana J, et al. Cytotoxicity of folpet fungicide on human bronchial epithelial cells[J]. Toxicology, 2008, 249(2-3): 160-166.

[4] Gudi R, Krsmanovic L. Nuclear aberration test in the mouse duodenum(folpet) and photomicrographs [R]. Rockville, MD: BioReliance Corporation, 2001. Report no. AA31S K. 123005.

[5] European Food Safety Authority. Peer review of the pesticide risk assessment of the active substance folpet[J]. EFSA Scientific Report, 2009, 297: 1-80.

[6] Tamano S, Kurata Y, Shibata M. 13-Week oral toxicity study of captafol in F344/DuCrj rats[J]. Fundam Appl Toxicol, 1991, 17(2): 390-398.

[7] Jeff SM, Jerome HG. Acetaminophen-induced forestomach lesion in normal rats following intravenous exposure[J]. Toxicol Pathol, 2011, 39(5): 861-866.

[8] Yoshihide U, Masaru T, Yasufumi O, et al. Gastric mucosal changes induced by polyethylene glycol 400 administered by gavage in rats[J]. J Toxicol Sci, 2011, 36(6): 811-815.

[9] Tomoyuki S, Masao H, Kazuo H, et al. Sequential morphological and biological changes in the glandular stomach induced by oral administration of catechol to male F344 rats[J]. Toxicol Pathol, 1999, 27(4): 448-455.

[10] Nyska A, Waner T, Paster Z, et al. Induction of gastrointestinal tumors in mice fed the fungicide folpet: possible mechanisms[J]. Jpn J Cancer Res, 1990, 81(6-7): 545-549.

[11] Iturri SJ, Soto J. Inhibitory Effect of folpet in the small intestine absorption capacity of the mouse [J]. Bull Environ Contam Toxicol, 1994, 53: 648-654

本刊中的类似文章

1. 李百祥, 张晓峰, 马若波. 小鼠前胃癌诱导中Cx32表达的变化[J]. 中国公共卫生, 2003, 19(9): 1057-1058
2. 柳云恩, 施琳, 张玉彪, 张继存, 侯明晓. 95%灭菌丹原药大鼠经口毒性实验结果评价[J]. 中国公共卫生, 0, (0): 0-0

文章评论 (请注意: 本站实行文责自负, 请不要发表与学术无关的内容! 评论内容不代表本站观点.)

反馈人	<input type="text"/>	邮箱地址	<input type="text"/>
反馈标题	<input type="text"/>	验证码	<input type="text" value="8496"/>