REVIEW ARTICLE

Occupational Health And Safety In Drug Manufacturing Industry: A Short Review

Muhammad Maaz Arif¹

1. Department of Medical Education, University of Health Sciences, Lahore, Pakistan **Corresponding author:**<u>maazarifbutt@gmail.com</u>

ABSTRACT

Background: Not much is known about the health risks of workplace in the drug manufacturing industry. While the pharmaceutical industries seem safe superficially, the production of medicines demand a well maintained and clean working atmosphere and the typical depiction of employees wearing white coats add to the illusion. Objective: To document occupational health and safety in drug manufacturing industry. Methods: Large-scale research was conducted to assess the health, safety and efficacy of specific drugs before their marketing and distribution, however only a few surveys have examined the occupational health and safety of employees who manufacture these drugs. **Results:** There are many irritants and drugs that cause risk to health, these include powdered penicillin, nicotinic compounds, Local anaesthetics, chloroform, sulfonechloramides, benzene, mepacrine, acriflavine and sulfonating agents. The most crucial aspects of efficient pharmaceutical safety management are knowing and recognising the danger of hazardous facility and hazard discharge. The easiest method to avoid difficulties is to eliminate the danger or employ substitution (choosing the least toxic or safest product or process available). Furthermore, it appears plausible to anticipate that chemical-related health impacts of pharmaceutical industry activity, long-term effects, in particular, are likely under-reported and understated. Other dangers (for example, ergonomic design and stressors) are reported more frequently in internal company systems, but are rarely acknowledged in public. Conclusion: General safety guidelines and rules for drug manufacturing safety should be followed. These include; not to work alone in the laboratory, never use pipette with mouth, wear goggles or safety glasses in the laboratory, practice personal hygiene rules, do not eat or drink in the laboratory, practice housekeeping use PPE and the of good techniques. Keywords: Occupational health and safety, drug manufacturing, Pharma industry, international standards

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INTRODUCTION

There is not much information about the health dangers associated with the drug manufacturing industry's workplace. While the pharmaceutical industries seem safe superficially, the making of medicines demands a well-kept and clean working atmosphere and the typical depiction of employees wearing white coats add to the illusion.¹ Large-scale research is conducted

to assess the health, safety and efficacy of specific drugs before their marketing and distribution, however, only a few surveys have examined the occupational health and safety of employees who manufacture these drugs. While the health and safety of chemical workers who manufacture nonpharmaceutical chemicals has been studied extensively, different is the case with pharmaceutical production employees. The morbidity and mortality experience of pharmaceutical employees has been previously discussed in a few researches.^{2, 3} The drug industry manufactures therapeutic substances, both human and veterinary medications, pharmaceuticals, and associated items, in a growing concentrated network of large international firms and

Materials and Methods

Study design: Systematic review

Setting: Department of Medical Education, University of Health Sciences, Lahore, Pakistan

Duration: The study was done in 6 months

Inclusion criteria: Full length article published in English language meeting our search strategy were taken

Exclusion criteria: Copy right and articles required subscription and fee were excluded. **Search strategy:** The articles were selected using Pubmed and Google Scholar databases any where from year 1990 onwards. The keywords used for searching were "pharmaceutical", "industry", "workers", "health" and "safety".

RESULTS:

Common epidemiological findings of drug industry employees have been carried out in the United States (excluding Canada) and Europe. The findings depicted an increase in the prevalence of chronic bronchitis and hypertension among Serbian plant workers.⁵ Furthermore, it was observed that among Polish sulfonamide workers, changes to mouth and teeth including leukoplakia (thick white patches) were reported.⁶ In a report that showed any worker who had a full-time job from 1970-1996 and with assessed exposure to 9 of the more than 50 chemicals in the plant, lung cancer that could have occupational non-Hodgkin's links to lymphoma and lymphatic-hematopoietic tissue cancer in a small group (majority male) employees of the same U.S plant.⁷

subcontracting facilities. The sector's activities are divided into five basic categories: administration, manufacturing, marketing and sales, distribution, and research & development (R & D). There are three essential processes in the production of drugs: R & D, manufacturing, and packaging.⁴

Another study reported in a Croatian plant displayed the danger of chronic and acute respiratory symptoms from various unnamed drugs.⁸

Chemical Hygiene Programme and a designated Chemical Hygiene Officer are required by the "Laboratory Standard," 29 CFR 1910.1450. The fundamental principles of safe laboratory work can be found in your local Chemical Hygiene Programme. The people responsible for the chemical hygiene program include the chief executive officer, hygiene officer. chemical laboratory supervisors and laboratory workers.⁹ It is vitally important to identify and control chemical hazards. Identifying chemical hazards include chemical labeling, the manufacturer, importer and precautionary statements.¹⁰ The control of chemical hazards includes elimination, substitution, engineering measures, isolation. administrative controls and personal protective equipment.¹¹

Major dangers are a concern for pharmaceutical facilities and others. Risk management firms' assistance in this area for pharmaceutical organizations is based on their experience working across the entire spectrum of these businesses. The method of managing major hazard risk is based on risk assessment in safety management, which comprises formal safetv management systems behavior-based and safety programs. Risk evaluation of laboratory tasks biological containment laboratories is required for diagnostic and detection

activities, as well as research and development. In these laboratories, many different types of microorganisms are treated, and each activity done on the microorganism has the potential to represent a risk.¹² As a result, the risks must be assessed so that precautions can be taken to

protect both employees and the environment from infection. Risk assessments, according to the WHO's laboratory biosafety manual, are the foundation of any laboratory safety program.¹³ Some many irritants and drugs cause risks to health, these are shown in Table 1.^{14, 15}

DRUG	HARMFUL EFFECTS
Powdered penicillin	Skin sensitizer and skin irritant
Nicotinic compounds	Itching, Diffused erythema
Local anesthetics	Epigastric pain, GIT problems, Liver diseases, Fatigue
Chloroform	GIT problems, Liver poison
2-methyl-1, 4-naphthoquinone	Skin sensitization, Dermatitis
Sulfonechloramides	Local irritants and sensitizers
Benzene	Benzene poisoning
Mepacrine and acriflavine	Conjunctivitis
Sulfonating agents	Irritant, Bronchitis, Conjunctivitis.

TABLE 1: Drugs with their harmful aspects ^{14, 15}

CONCLUSION AND RECOMMENDATIONS:

The most crucial aspects of efficient pharmaceutical safety management are knowing and recognizing the danger of facilities hazardous and hazardous discharge. An effective hazard and risk assessment enables the development of an incident action plan as well as the implementation of strategies and tactics. A precautionary attitude to hazards, as in other work environments, is the basis of a safe and healthy pharmaceutical industry workplace, and responses to dangers should be guided by public health principles. The easiest method to avoid difficulties is to eliminate the danger or employ substitution (selecting the least toxic or safest product available). Furthermore, it appears plausible to anticipate that chemical-related health impacts of pharmaceutical industry activity,

particularly chronic effects may be under-

reported orincorrectly estimated. Other dangers (for example, ergonomic design and stresses) are recorded more frequently in internal business processes but are rarely highlighted in the mainstream media. General safety guidelines and medication production safety rules must be observed. These include not working in the laboratory alone. Never use a pipette in your mouth. Always wear goggles or safety glasses in the laboratory. Follow personal hygiene guidelines (for example, wash your hands before exiting the laboratory). In the laboratory, no food or drink should be permitted. Wear PPE andavoid wearing laboratory coats outside of the laboratory and close-toed, sturdy shoes. Use good housekeeping procedures, such as keeping paths free and labeling and dating all containers. Improving existing regulations'

ratification and implementation. Increasing the scope of risk assessments and controls. Multi-agency approaches to chemical management are being improved and raised awareness. It is critical to pay close attention to the criteria for reproductive and genotoxic compounds, as well as mutagens and carcinogens. Ergonomic requirements must also be met.

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