

International Journal for Pharmaceutical Research Scholars (IJPRS)



ISSN No: 2277 - 7873

# **RESEARCH ARTICLE**

# Microbial Assessment of Some Common Indian Brands of Talcum Powder

Wahla V\*, Kasana M

Department of Microbiology, Kanya Gurukul Girl's Campus, Gurukul Kangri University, Jwalapur, Haridwar, 249407 Uttarakhand, India.

Manuscript No: IJPRS/V4/I2/00102, Received On: 19/05/2015, Accepted On: 30/05/2015

#### ABSTRACT

A total of three samples of talcum powders were examined for their total aerobic bacterial counts. A serial dilution technique was carried out and plating was done by using standard spread plate technique. The bacterial load of talcum powder Boroplus ranges from  $1.5 \times 10^7$  to  $2.1 \times 10^8$  cfu/g with a mean bacterial load of  $1.8 \times 10^8$  cfu/g, in Dermicool ranges were  $1.6 \times 10^9$  to  $2.8 \times 10^7$  cfu/g with a mean bacterial load of  $2.3 \times 10^8$  cfu/g and in talcum powder Navratna colonies were uncountable. Bacteria isolated from talcum powder were *Staphylococcus* spp. 40% and *Bacillus* spp. 60%. In conclusion, the talcum powders studied showed to be more heavily contaminated. This may be as a result of poor manufacturing process, poor hygiene and contaminated raw materials.

#### **KEYWORDS**

Talcum powder, Bacterial load, Bacillus and Staphylococcus

## INTRODUCTION

Talcum powder is a cosmetic product used by both men and women to improve their looks and avoid the growth of bacterial pathogen which may cause unpleasant odor and sometimes skin infections. One of its most common uses is in baby care, to reduce irritation from diapers. The field of cosmetics and microbiology had not come into contact much before the 1930s and cosmetic microbiology became more important in 1940s.<sup>1</sup> Talcum powders have positive effects on adult and babies skin, which mainly used to smooth the appearance of skin but negative effects also occur if they were contaminated. It was reported that some of these talcum powders are contaminated with spores of microorganism and can support their growth when they are poorly preserved and causes several disease.<sup>2</sup>

\*Address for Correspondence: Verinder Wahla Department of Microbiology, Kanya Gurukul Girl's Campus, Gurukul Kangri University, Jwalapur, Haridwar, 249407, Uttarakhand, India. E-Mail Id: verinderwahla19@gmail.com The warm and rather humid climatic conditions would tend to support the survival and growth of many microorganism, so rapid growth and multiplication would be expected. This could lead to biodegradation of the product and hence the risk of microbial contamination to consumers of the product.<sup>3</sup>

In 2014 Omorodian and his coworker isolated bacteria from the baby powder were *Staphylococcus* spp., *Bacillus* spp., *Streptococcus* spp., *Micrococcus* spp., and *Escherichia coli*, while bacterial isolates from adult powder were *Staphylococcus* spp., *Bacillus* spp., *Streptococcus* spp., and *Micrococcus* spp.

*Escherichia coli* were not isolated from any of the adult powders. This possibly is due to over confidence in the traditionally good hygienic conditions under which such products are manufactured and also because it is assumed that added preservative will prevent microbial growth upon storage and during use.<sup>4</sup> The objective of this study is to assess the microbial quality of some selected brands of commonly used adult powder to recommend the possibility of some health risk to consumers. To overcome from this problem the packaging of talcum powder must be done in aseptic condition and dry places, and machinery used for grinding must be free from microbial contamination and to avoid the skin problem caused by the contaminated talcum powder, antibiotic must be used. Antibiotic are used to fight against bacterial infections. Kirby-Bauer antibiotic testing (KB testing or disc diffusion antibiotic sensitivity testing) is a test which uses antibioticimpregnated wafers to test whether bacteria are affected by antibiotics.<sup>5</sup>

#### MATERIAL AND METHODS

#### **Sample Collection**

A total of 3 commercial sample of talcum powder brand name Navratna, Boroplus and Dermicool were purchased from shop of Roorkee, Haridwar (Uttarakhand). For precaution, check the seal of product and transported to the laboratory and analyzed.

#### Bacteriological Counts of the Cosmetic Powders

A stock solution was prepared by dissolving 1gm

of sample into 9 ml of sterile distilled water. A tenfold serial dilution was made and last three dilutions transferred into NAM plates using spread plate method. The plates were allowed to solidify and incubated at  $37^{0}$ C for 24-48 h.

#### **Identification of Bacterial Isolates**

All bacterial isolates were identified based on their Gram reaction and biochemical reactions (Table 2).

#### RESULTS

The results obtained shows that the bacterial load of talcum powder Boroplus ranges from  $1.5 \times 10^7$  to  $2.1 \times 10^8$  cfu/g with a mean bacterial load of  $1.8 \times 10^8$  cfu/g.

The bacterial load of talcum powder Dermicool were ranges from  $1.6 \times 10^9$  to  $2.8 \times 10^7$  cfu/g with a mean bacterial load  $2.3 \times 10^8$  cfu/g and in talcum powder Navratna colonies were uncountable (Table 1). Out of 3 samples of talcum powders that were analysed, bacteria isolated from Boroplus powder were *Staphylococcus* spp. 15%, *Bacillus* spp. 85% while bacterial isolates from Navratna powder were *Bacillus* spp. 100%, *Staphylococcus* spp. were not isolated from this powder and bacterial isolates from Dermicool powder *Staphylococcus* spp. 40% and *Bacillus* spp. 60%.

S.No	Sample	Dilution	Dilution factor	No. of colonies	C.F.U./g	Mean ± SD
1.	Boroplus	10 <sup>-4</sup> 10 <sup>-5</sup> 10 <sup>-6</sup>	$10^4 \\ 10^5 \\ 10^6$	154 216 172	1.5 X10 <sup>7</sup> 2.1 X10 <sup>8</sup> 1.7 x 10 <sup>9</sup>	1.8±3.18x10 <sup>9</sup>
2.	Navratna	10 <sup>-4</sup> 10 <sup>-5</sup> 10 <sup>-6</sup>	$10^4 \\ 10^5 \\ 10^6$	TNTC TNTC TNTC	TNTC TNTC TNTC	TNTC
3.	Dermicool	10 <sup>-4</sup> 10 <sup>-5</sup> 10 <sup>-6</sup>	$10^4 \\ 10^5 \\ 10^6$	280 252 162	2.8X10 <sup>7</sup> 2.5X10 <sup>8</sup> 1.6X10 <sup>9</sup>	2.3±6.16x10 <sup>9</sup>

 Table 1: Enumeration of bacteria from talcum powder

 $\overline{\text{TNTC}} = \text{Too numerous to count}$ 

Table 2: Morphological and biochemical characteristics of bacterial isolates from talcum powder

	<b>Bacterial Isolates</b>		
Characteristics	B1	B2	
Grams staining	Gram	Gram	
Colony	White, irregular margin	Yellow, smooth margin	
Cell shape	Rod	Cocci	
Cell arrangement	Single	In cluster	
Catalase	+ve	+ve	
Lactose	-ve	A	
Sucrose	А	А	
Dextrose	А	Α	
Starch hydrolysis	+ve	-ve	
Nitrate reduction	-ve	+ve	
MR	-ve	+ve	
VP	-ve	-ve	

Table 3: Showing the order of drug susceptibilityin Bacillus spp.

Antibiotic	Content (mcg)	Zone of inhibition (mm)
Ampicillin (AS)	20	9
Co-Trimoxazole (BA)	25	2
Cephalexin (PR)	30	15
Tetracycline (TE)	30	25
Cefotaxime (CF)	30	10
Ciprofloxacin (RC)	5	31
Levofloxacin (QB)	5	24
Linezolid (LZ)	30	28
Cloxacillin (CX)	1	1
Roxythromycin (AT)	15	R
Lincomycin (LM)	2	4
Gentamicin (GM)	10	3

R = Resistance

Table 4: Showing the order of drug susceptibilityin Staphylococcus spp.

Antibiotic	Content (mcg)	Zone of inhibition (mm)
Amoxycillin (AM)	10	R
Cefazolin (CF)	30	R
Cephalexin (CP)	30	R
Roxythromycin (TH)	30	R

-ve=Negative

+ve=Positive

A=Acid

## Antibiotic Test

Antibiotic sensitivity study of different antibiotics on *Bacillus* spp. and *Staphylococcus* spp. shown result that *Bacillus* sensitive for all antibiotics of multidisc but resistant for antibiotic Roxythromycin and *Staphylococcus* spp. was sensitive only for antibiotic Erythromycin, while resistant for all other antibiotics of multidisc.

Cefadroxil (CD)	30	R
Erythromycin (E)	15	19
Ciprofloxacin (CL)	5	R
Vancomycin (Vn)	30	R
Ofloxacin (OF)	5	R
Sparfloxacin (SP)	5	R
Ampicillin (I)	10	R
Cloxacillin (V)	5	R

## R = Resistance

## DISCUSSION

Based on the findings of this work, the talcum powders analyzed were contaminated with Gram positive bacteria. In a similar study, Hugbo et al.,  $(2003)^6$  isolated *Staphylococcus spp.* and other Gram positive cocci were the most predominant, Gram negative isolates were hardly found Nasser  $(2008)^7$  also reported more of bacterial than fungal contamination. Dashen *et al.*,  $2011^2$ isolated Staphylococcus aureus, Clostridium tetani, Bacillus spp. and (Ashour et al., 1989)<sup>8</sup> isolated Escherichia Enterobacter coli. agglomerans, **Staphlococcus** aureus. and Citrobacter freundii.

The International Microbiological standard recommended limit for bacteria contaminants in cosmetic products is  $1.0 \times 10^3$  CFU/g for bacteria. It was observed in our study the total bacterial counts were above the recommended limits. High microbial quality was observed in this study could be caused by poor manufacturing practice and improper storage. Bacterial isolates from adult powder were *Staphylococcus* spp. 48%, *Bacillus* spp. 6%<sup>3</sup> but from talcum powders in our study we found two dominating bacteria

*Staphylococcus* spp. and *Bacillus* spp. In present study we found 40% *Staphylococcus* spp. and 60% *Bacillus* spp.

The frequency of occurrence of bacteria in the total sample shows that all the samples are contaminated with bacteria which indicating that talcum powders can permit the growth of bacteria. It was also observed that Gram positive organisms were the predominant contaminants in the powders. The high bacterial counts obtained may be due to poor storage, manufacturing handling. Bacillus practice or and Staphylococcus spp. in talcum powder causes skin irritation. Generally, the results obtained from this study showed that these talcum powder were highly contaminated.<sup>3</sup>

# CONCLUSION

The conclusion drawn from this study shows that all the three samples of talcum powder have capable of causing health hazards due to high microbial loads. The microbial loads are above standard. This may be as a result of poor manufacturing practices, poor hygiene, contaminated raw materials or the susceptibility of the ingredients contained in the talcum powders. The presence of organisms such as *Bacillus* spp. and *Staphylococcus* spp. in the talcum powders implies that they can serve as vehicles for the transmission of disease.

## REFERENCES

- Gamal, M. A., Abo Azza, M. M., & Al Gayeed, A. O. Microbiological Quality Assessment of Some Brands of Cosmetic Creams Sold Within Alkhoms City, Libya.
- Dashen, M. M., Chollom, P. F., Okechalu, J. N., & Ma'aji, J. A. (2011). Microbiological quality assessment of some brands of cosmetics powders sold within Jos Metropolis, Plateau State. *Journal of Microbiology and Biotechnology Research*, 101-106.
- Omorodian,, Nnenna, J. P., Ezediokpu, M. N.; Edward, G. (2014). Microbiological quality assessment of some brands of cosmetics powders sold within port Harcourt rivers state, *Nigeria*. 7-11.

- Raghad, A. R.; Ebtihal, N. S. and Heyan, H. (2010). A study of Microbiological contamination in cosmetics and toiletries in Iraq. Contamination of talcum powder and body lotion.
- Bauer, A. W., Kirby, W. M. M., Sherris, J. C. T., & Turck, M. (1966). Antibiotic susceptibility testing by a standardized single disk method. *American Journal of Clinical Pathology*, 45(4), 493.
- Hugbo, P. G., Onyekweli, A. O., & Igwe, I. (2005). Microbial contamination and preservative capacity of some brands of cosmetic creams. *Tropical Journal of Pharmaceutical Research*, 2(2), 229-234.
- 7. Nasser, L. A. (2008). Fungal profiles isolated from open and used cosmetic

products collected from different localities in Saudi Arabia. 15 (1), 121-128.

- 8. Ashour, M. S. E., Abdelaziz, A. A., Hefni, H., & El-Tayeb, O. M. (1989). Microbial contamination of cosmetics and personal care items in Egypt—body lotions and talcum powders. *Journal of Clinical Pharmacy and Therapeutics*, *14*(3), 207-212.
- 9. Fujital, P. G. and Onyeard, A. (2005). Microbial contaqmination and preservation capacity of some brands of cosmetic creams. *Tropical Journal of Pharmaceutical Research*, 2: 229-234.
- Razooki, R., Saeed, E., & Hamza, H. (2009). A study on Cosmetics Products Marketed in Iraq: Microbiological Aspects. *Iraqi Journal* of Pharmaceutical Sciences, 18(2), 20-25.

