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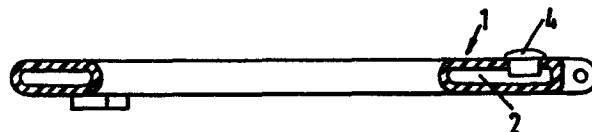
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⑤④ **Lavatory seat.**

⑤⑦ The invention provides a plastics lavatory seat (1) which has an internal channel or cavity (2) for containing liquid disinfectant and/or deodorant (3) of the phenolic or cationic kind, the liquid (3) and the plastics being compatible and the plastics being permeable to and capable of absorbing the liquid so that a hygienic seat is obtained.



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"Lavatory Seat"

The invention relates to a lavatory seat.

A lavatory seat is known in which vapour of a disinfectant and/or deodorant is supplied to a
5 surface of the seat through pores through the thickness of the material of the seat, (United Kingdom Patent No. 1 340 075). In this prior art specification the pores extend between the surface and a hollow interior of the seat in which liquid disinfectant and/or deodorant is housed. However,
10 although it has been found that this seat is hygienic on its upper surface there is sometimes not such a good bactericidal action in the underside of the seat and it is on the underside that there is sometimes spread of infection by splashing from a
15 lavatory bowl to which the seat is affixed.

The invention as claimed is intended to provide a remedy. It solves the problem of how to provide a lavatory seat in which bacteria are substantially
20 eliminated, thereby providing a seat which is hygienic over all its surfaces.

The advantages offered by the invention are mainly that the seat, being formed of plastics material which is permeable to liquid disinfectant,
25 can absorb disinfectant so that the whole seat is substantially bacteria free, while maintaining body supporting surfaces of the seat "dry". Also the

plastics is coherent, not having pores, and is easily manufactured as by moulding and provision for providing pores does not have to be made.

5 The material, which may comprise a body supporting surface of the seat and a boundary surface for part of the cavity, may be a plastics material.

The plastics may comprise perspex, polypropylene, acrylonitrile butadiene styrene (ABS), polyvinyl chloride or polyethylene.

10 The whole seat may comprise the plastics.

Where the whole seat is made of plastics, it may be formed by blow moulding, in for example a one-shot moulding process.

15 The seat may include means for charging the cavity with disinfectant and/or deoderant. The charging means may comprise a removable filler cap.

The filler cap may be transparent and may be so arranged that there is in use an air gap between its lower surface and the upper level of liquid in the (horizontal) seat. The air gap ensures that
20 when the seat is raised and lowered, the (liquid) disinfectant and/or deodorant, in liquid form, flows around the cavity which ensures a good mixing of the liquid, while the transparent nature of the
25 cap allows the colour of the liquid to be monitored. This is to enable the state of the disinfectant to be controlled, because it changes colour as it becomes spent.

30 There may be means to secure the cap against tampering e.g. by vandals.

One way of carrying out the invention is described in detail below, with reference to drawings which show only one embodiment, in which:-

35 Figure 1 is a plan view of a lavatory seat in accordance with the invention;

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Figure 2 is a longitudinal sectional view of the seat of Figure 2; and

Figure 3 is an enlarged sectional view of part of the seat of Figures 1 and 2, at the filler cap.

5 The Figures show a lavatory seat 1 which has a liquid disinfectant distributing device in the form of an internal cavity 2 which extends throughout the whole seat. In accordance with the invention, the cavity 2 is for containing liquid disinfectant
10 2 (Figure 3) (though liquid deodorant could be used alternatively or in addition). The cavity 2 is wide but not deep and is made integrally with the seat 1 when that seat is made by blow moulding polyvinyl chloride. The polyvinyl chloride is permeable to
15 the disinfectant 3. The cavity 2 is filled through charging means in the form of a filler cap 4. The cap 4 is transparent and is so constructed and arranged that there is an air gap 5 between it and the inner surface 6 of the upper boundary of the
20 cavity 2.

 In accordance with a preferred embodiment of the invention and in use, the cavity 2 is filled with water through the charging opening when the cap 4 is removed. Disinfectant in liquid capsule,
25 tablet, powder or paste form is then added to the water and disperses or dissolves in it. The cap is placed in position in the seat.

 When the seat 1 is raised and lowered, the liquid disinfectant flows round the cavity and is
30 thoroughly mixed and also thoroughly contacts the material of the seat. This movement is provided for by the air gap 5, into and out of which the liquid can flow.

 The disinfectant slowly permeates through the
35 material of the seat, so disinfecting all its surfaces

and rendering them hygienic, but it does not "wet" the seat which is therefore comfortable to use. The rate of penetration depends on the thickness of the plastics used. It will also be understood that
5 the water is a carrier for the disinfectant which it brings into intimate contact with the seat and enhances its penetration therethrough.

The disinfectant has a particular colour. As it becomes spent, its colour changes. This change
10 can be observed through the transparent cap 4 and, when required, more disinfectant, paste or the like can be added to re-charge disinfectant to restore it to its correct strength. Of course, the seat can be made in other ways, for example by moulding,
15 injection moulding, rotational moulding, a plastics spinning operation or in any other suitable way such as cellular blow moulding of a suitable expandible plastics. In this case the body of the seat would comprise a thin skin backed by a porous or foam
20 structure, but there would be no pores extending through the skin. The open pore structure would facilitate the diffusion of the disinfectant.

The disinfectant may be a phenolic disinfectant such as:-

- 25 (a) Chloroxlenol (4-chloro-3:5 -xylenol)
(b) Chlorocresol (4-chloro -3 -methylphenol)
(c) Sudol (a proprietary blend of a closely cut fraction of phenols, chiefly xylenols and ethyl phenols.)

30 The phenolic system contains 50% phenols solubilised by vegetable soap.

Alternatively the disinfectant may be a cationic disinfectant such as chlorhexidine gluconate (1:6 -di -CN -4 -chlorophenyl -di -guanido) -hexane
35 digluconate).

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Chlorhexidine gluconate is totally miscible with water and Sudol is formulated in a vehicle which allows ready dilution with water. Chlorocresol and chloroxylenol are both sparingly soluble in water (0.4% and 0.03% respectively) and can be dissolved in aqueous polyethylene glycol (PEG 400) to form cosolvent mixtures. PEG 400 is an acceptable cosolvent for external medicinal use as on a lavatory seat.

It has been found that the total uptake, and hence the "reservoir" of disinfectant in the plastics material of the seat is greater for aqueous chlorocresol, than for chloroxylenol in 10% PEG 400.

Plastics material which can be used are, in addition to the unplasticised PVC mentioned, filled acrylonitrile butadiene styrene (ABS), cellulose-acetate butyrate (CAB), polypropylene and polyurethane. I have found that the permeation rate of chlorocresol through CAB is ten times greater than through polypropylene, while still maintaining the plastics in a "dry" state, that is to say the plastics was not uncomfortable to use.

In both cases, too, there was considerable sorption of the disinfectant by the plastics, so that the whole body of plastics comprising the seat was rendered hygienic and disinfected.

The efficacy of plastics permeable to liquid disinfectant for removing organisms or bacteria, is shown in the following specific Example.

EXAMPLE

ORGANISMS

<u>Escherichia coli</u>	NCTC 8196
<u>Staphylococcus aureus</u>	NCTC 6571
<u>Pseudomonas aeruginosa</u>	NCTC 6749
<u>Streptococcus faecalis</u>	NTCT 775

All organisms were grown to stationary phase in nutrient broth, filtered and resuspended in non nutritive buffer at a concentration of approximately 5×10^7 orgs. ml⁻¹

5 PLASTICS

The plastics is polypropylene. Discs of 2.5 cm diameter were cut out and equilibrated with the disinfectant chlorocresol.

INOCULATION LEVELS

10 20 µl of suspension in salt solution with or without 10% serum containing 5×10^6 organisms ml⁻¹ were inoculated onto each disc as approximately 10 x 2 µl droplets. This gives 10^5 organisms per disc which is approximately 2×10^4 organisms cm⁻².

15 (This is in the high end of the range of contamination levels encountered on the surface of hospital toilet seats.)

PROCEDURE

20 Inoculated discs were stored at 20°C in glass vessels adjusted to a range of humidities by the use of saturated salt solutions. After various time intervals discs were removed and washed in salt solution. The number of viable bacteria in the suspensions obtained was assessed by serial dilution
25 and plating on nutrient agar. Survival, expressed as a fraction of the number inoculated, was calculated by counting the colonies formed after 48 hours at 37°C.

RESULTS

30 1. The sensitivity of the four organisms to the antimicrobial agent chlorocresol as assessed by "Minimum Inhibitory Concentration" tests showed that E. coli and Staph. aureus were inhibited by 0.25% chlorocresol and Pseudomonas aeruginosa and Strep-
35 tococcus faecalis were inhibited by 0.5%.

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2. The survival of the four organisms when exposed to drying at room temperature and ambient relative humidity indicated that the gram negative organisms, E. coli and Ps. aeruginosa behaved in a similar fashion.

It will be understood that the lavatory seat illustrated in the drawings and above described can be modified in various ways. Thus the cavity 2 may not be necessary if a suitable polar polymer such as nylon were used because the cellular structure provides the route for the transport of the disinfectant to the surface of the seat. Also the skin or surface layer could be of a different material to the (foamed) core. If the seat is of foamed plastics without a cavity 2, there may be a feed channel for distribution liquid disinfectant throughout the seat. If the channel is a surface groove, a permeable or microporous sheath may be insertable over the seat to cover the groove and through which the disinfectant can pass.

Claims:

1. A lavatory seat characterised in that it is made of a material which is permeable to a liquid disinfectant and/or deodorant.
2. A lavatory seat according to Claim 1, characterised by a device (2) for distributing disinfectant (3) through the liquid disinfectant permeable material of the seat (1).
3. A lavatory seat according to Claim 1 or Claim 2, characterised in that the liquid disinfectant permeable material of the seat (1) is a plastics material.
4. A lavatory seat according to Claim 3, characterised in that the plastics material of the seat (1) is polypropylene, acrylonitrile butadiene styrene, polyvinyl chloride or polyethylene.
5. A lavatory seat according to Claim 4, characterised by liquid disinfectant and/or deodorant (3) in the distribution device (2).
6. A lavatory seat according to Claim 6, characterised in that the liquid (3) is a phenolic or a cationic disinfectant.
7. A lavatory seat according to any preceding claim, characterised by a filler means (4).

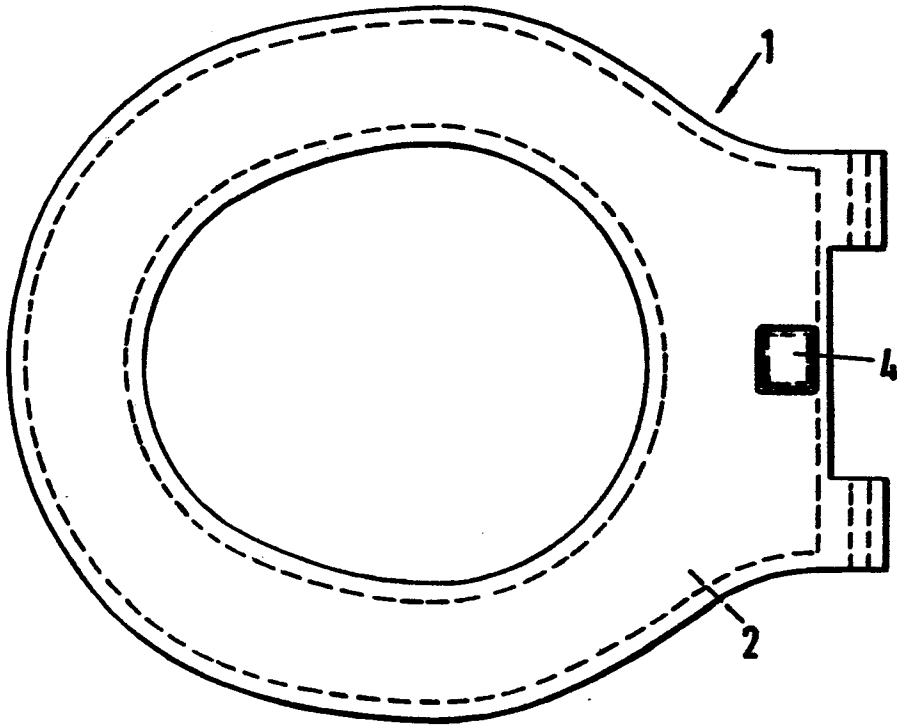


FIG. 1.

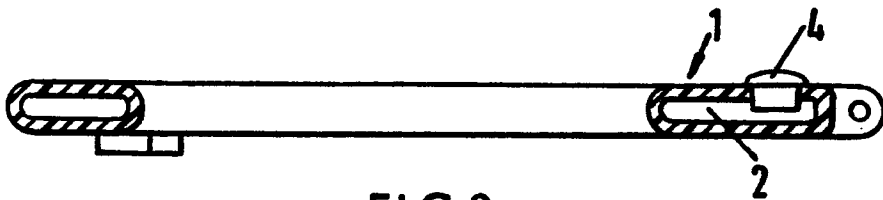


FIG. 2.

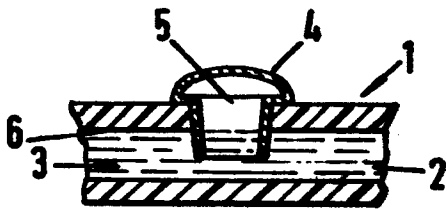


FIG. 3.



DOCUMENTS CONSIDERED TO BE RELEVANT			CLASSIFICATION OF THE APPLICATION (Int. Cl.)
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	
X	<u>DE - C - 69 827 (ROSE)</u> * The whole document * ---	1,2,5,7	A 47 K 13/30
	<u>GB - A - 331 157 (CLIFFORD)</u> * The whole document * ---	3,6	
	HUGO W.B.(ED.) "Inhibition and destruction of the microbial cell", 1971, Academic Press, London * Pages 95-96, 549, 438-439 * -----	6	
			TECHNICAL FIELDS SEARCHED (Int.Cl.) A 47 K 13/00 A 47 K 13/02 A 47 K 17/00 A 47 K 13/24 A 47 K 13/30
			CATEGORY OF CITED DOCUMENTS X: particularly relevant A: technological background O: non-written disclosure P: intermediate document T: theory or principle underlying the invention E: conflicting application D: document cited in the application L: citation for other reasons
The present search report has been drawn up for all claims			&: member of the same patent family, corresponding document
Place of search	Date of completion of the search	Examiner	
The Hague	03-10-1978	BAERT	