Is chronic detergent ingestion harmful to the gut?

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ABSTRACT Synthetic detergents are used in large quantities as household and industrial cleaners. Because of the common practice of leaving dishes washed in detergent solutions to dry without rinsing these compounds are ingested. We have calculated that an adult takes in about 1 mg/kg detergent a day and babies can be administered between seven and 10 mg/kg a day. Rats were fed a dose of 100 mg/kg a day in a pilot experiment and gross abnormalities were found in the gastrointestinal tract, the most striking being subtotal villous atrophy of the small bowel mucosa and glandular atrophy in the colon. These changes were not reversible 12 weeks after cessation of detergent administration.

The development of synthetic detergents 30 years ago revolutionised cleaning processes. These comparatively cheap materials are used in the manufacture of washing up liquids, bath oils and foams, shampoos, soap powders, and numerous other household and industrial cleaners. It is common practice to leave glasses, crockery, and eating implements to dry with a coating of detergent, and indeed some manufacturers recommend this since the fluorescent additives cause them to “sparkle.” These compounds are inevitably ingested from such implements. Feeding bottles for babies are often steeped in strong solutions of detergent and not rinsed between feeds.

Detergents have a potent effect on cell membranes. At low concentration they form complexes with brush border enzymes and at higher concentrations solubilise membranes by forming micelles with the lipid protein components. They have provided membrane biologists with a useful tool for the study of membrane components in vitro. What are the effects of these substances in vivo? We present a descriptive account of the effects of chronic ingestion of synthetic detergent on the morphology of the gastrointestinal tract of the rat.

Materials and methods

Six female Wistar rats, litter mates of average weight 200 g, were caged together and fed a standard 41B diet. The drinking water was a 1% solution of a household detergent (Fairy Liquid: Proctor & Gamble 37/8077/G) in tap water.

The animals were weighed regularly and the faeces inspected daily. The amount of solution consumed each day was noted. Single, randomly chosen rats were killed at regular intervals between 18 and 68 weeks by ether inhalation in a closed jar, and postmortem examination was performed immediately. After inspection of the viscera in situ, a 6 Ch polyethylene cannula was passed into the stomach through the mouth and the entire gastrointestinal tract lavaged gently with physiological saline, thus clearing it of intestinal contents. The bowel was then filled with 10% buffered formol saline. Segments of intestine were then isolated and tied so as to retain intraluminal formalin and to ensure immediate mucosal fixation. These segments comprised oesophagus to duodenojejunal junction, small bowel to just proximal to the ileocecal junction, and finally caecum to rectum. They were placed in 10% buffered formol saline for external fixation. All accessory organs of digestion were excised and fixed for 24 hours.

After fixation, the gastrointestinal segments were cut open along the antimesenteric border and the mucosa inspected. Segments were then made into a “swiss roll” and processed for histological examination. This technique allows microscopic examination of the mucosa of the entire gastrointestinal tract. Processed tissues were embedded in wax and cut at 4u on a Leitz rotary microtome before staining with haemotoxylin and eosin, PAS, and alcian blue.

Fairy Liquid consists of 40% w/v anionic alkyl benzene sulphonate, approximately 60% w/v nonionic alkyl ether sulphate, and a small amount of blue-green colour agent. One of the components of
this detergent possesses an absorption peak at 235
mm and was used to quantify the amount of
detergent remaining on an unrinsed standard 23 cm
diameter dinner plate, a 230 ml glass tumbler, and a
baby’s feeding bottle. A washing solution of 5 ml
detergent in 2 l of tap water was made (this is prob-
able less than the concentration a housewife would
use for dishwashing). Dilutions of this solution were
read in a Unicam SP 1700/1800 ultraviolet spec-
trophotometer (Pye Unicam Ltd) at 235 nm, using
tap water as a zero reference, and optical density
was plotted against concentration. The utensils were
washed in the fluid and allowed to dry. The recepta-
cle surface of each utensil was washed with 20 ml of
water and this fluid read in the spectrophotometer.
An additional quantification was made for the
baby’s feeding bottle after steeping in a strong solu-
tion of detergent solution (10 ml/l).

Results

The dilution curve of optical density against con-
centration obeyed Beer’s Law and gave a straight
line through the axis. From this plot the concentra-
tion of detergent remaining on the utensils was cal-
culated, assuming even dispersion of fluorescent
detergent molecules in solution. For three meals, six
cups of coffee, tea, or water, and a glass of beer the
amount of detergent ingested a day was calculated at
75 mg. Calculation for a standard feeding bottle at
two detergent concentrations showed that a baby
can be administered dosages of anything between
150 and 250 mg of detergent daily. Each rat con-
sumed 35 mg detergent a day.

The animals gained weight steadily. The growth
curves of the detergent fed rats were compared with
a mean growth curve of 10 control female Wistar
rats of initially similar age and weight over the same
period. There was little difference in the rate of
weight gain over the period of growth, but the max-
imum attained weight of the test rats was below that
of the controls. There were no external signs of ill-
ness, and none of the animals developed diarrhoea.

At necropsy the bowel wall of all the rats was
noted to be thinner and more transparent than nor-
mal. This became more obvious during luminal lav-
age.

Histology

The observed pathological changes although
remarkably constant in all the rats, tended to be
more florid in those killed later.

Tongue—There was gross hyperkeratinisation of
the surface of the tongue with total disappearance of
the taste buds in all rats.

Oesophagus—Large areas of the oesophagus were
denuded of mucosa but where present, this was thin
and friable with an uneven, frayed, and easily
detachable keratin layer. This attenuated mucosa
contained only a single layer of basal cells compared
with three or more in the normal rat oesophagus. No
strictures were seen, and there was little inflamma-
atory reaction in the ulcerated areas, but the features
were compatible with corrosive oesophagitis.

Stomach—The stomach appeared histologically
normal in all specimens examined.

Small intestine—Grossly abnormal changes were
seen throughout the small bowel. Villi were consid-
erably reduced both in number and size. They were
oedematous and friable and few were frankly necro-
tic. Extensive atrophy of the mucosal glands was
evident through cell hyperplasia. These changes
were progressively more noticeable in the distal part
of the small bowel. Mitotic figures were so few as to
suggest a failure of normal regenerative activity.
There was slight cellular infiltration in the lamina
propria. Submucosal lymphoid aggregates were
more prominent in rats killed at an early stage but
particularly sparse in those killed later (fig 1).

Large intestine—There was generalised glandular
atrophy and glands were small but widely separated
in the stroma. The swollen and distorted columnar
epithelial cells were seen sloughed off from the
mucosa (we are sure this is not autolytic surface loss
because of the method of preparation described

Fig 1 Longitudinal section of part of a roll of jejunum
showing sparse stunted villi and glands and numerous
goblet cells. (Haematoxylin and eosin × 100)
above). The slight cellular infiltration seen was confined to mucosa and submucosa without penetrating the muscularis (fig 2).

Liver—The liver showed an almost normal histological picture with mild periportal fatty infiltration.

Pancreas—Sections of the pancreas were difficult to cut because of microcalcification. These deposits tore gaps in the tissue and blunted the knife. Efforts to show this calcification radiologically in the anaesthetised rat and later in the pancreatic tissue itself, using low kilovoltage mammographic equipment, were not convincing, presumably because of the size of these deposits. The remaining pancreatic parenchyma appeared normal.

Discussion

There have been reports of corrosive oesophagitis after drinking dishwater detergent¹ and acute colitis after administration of detergent enemas.² The use of gastrograftin enemas in the treatment of meconium ileus³ resulted in several cases of fulminating enterocolitis with delayed perforation of the bowel, for which there was no rational explanation at the time. Gastrograftin contains the synthetic detergent Tween 80. The effect of detergent ingestion on lipid absorption has been reported, both after a single dose of hydrophobic detergent (138 g of Plurome L-81) in rats and after chronic feeding of a 1% solution of this detergent, over seven weeks.⁷ Abnormalities of fat absorption with steatorrhoea were described in these experiments.

We know that manufacturers test their products to a 1% concentration, but we are not aware of any published reports of their findings. We have shown profound morphological changes in the gastrointestinal tract in all our rats with this concentration.

Clearly, it would be inappropriate to extrapolate these results to man, particularly after a pilot experiment where our animals received dosages of, on average, 100 mg/kg a day compared with 1 mg/kg a day for an adult. Nevertheless, we would like to draw attention to the potential dosages of between 35 and 50 mg a day in babies weighing about 5 kg and think it possible that early mucosal damage may lead to chronic bowel disease.

Detergents are a useful if not indispensable part of modern living, but we should take precautions against eating them.

References