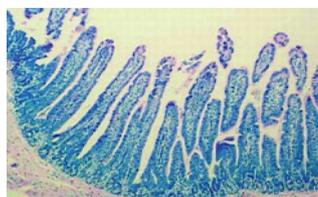


# The antibacterial effects of dietary zinc oxide in weaned piglets

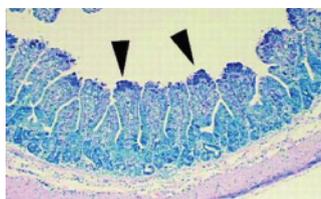
Dietary zinc oxide is known to increase growth and reduce post-weaning diarrhoea in piglets. But why do the other forms of zinc not perform as well as zinc oxide? Let's focus on the antimicrobial activity of zinc oxide.

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At weaning, when the intestinal microbiota changes and the immune system is not fully developed, pathogenic bacteria like enterotoxigenic *E. coli* (ETEC) find a favourable environment to multiply and colonise in the gastrointestinal tract of piglets. This manifests in the diarrhoea



**E. coli ileal adherence with zinc supplementation (above) and without (below).**



that we frequently observe post-weaning, combined with reduced growth.

Among all the additives available to the feed industry, zinc in the form of zinc oxide is a very popular solution. It is well established that zinc oxide exerts antimicrobial activity and recent studies with combined deep sequencing and PCR amplification offer new insights into the effect of the pharmacological dosage (3kg/T) on the intestinal microbiota of weaned piglets.

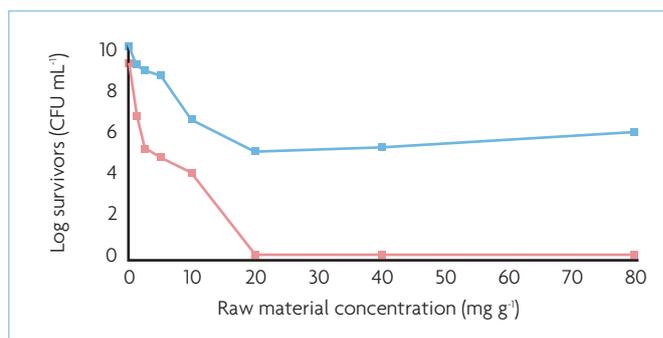
In vitro, the minimal inhibitory concentration (MIC) of ZnO against 75 strains from pig intestine showed that it is not possible to classify zinc sensitivity of bacteria according to their taxonomic origin.

However, strict anaerobic strains generally showed more diverse MIC than lactic acid bacteria or enterobacteria. Instead of a too simple separation between Gram+ and Gram-, the effect of zinc oxide seems to be species specific.

Under ex vivo conditions, it has been shown that high levels of ZnO leads to bacterial growth depression in the stomach and jejunum of weaned piglets, lactobacilli being more susceptible to high zinc dosage than other bacteria.

When piglets are fed 3000mg/kg zinc oxide, enterobacterial diversity increases, with reduced Enterobacteriaceae, Escherichia group and Lactobacillus species (see Fig. 1).

Even if we gained more understanding of the effect of ZnO on gut microbiota of weaned piglets, some questions remain:



**Fig. 2. Decrease of microbial populations with respect to zinc concentration for ZnO-1 suspension (red squares) or its supernatant (blue squares).**

- How does zinc oxide impair bacterial metabolism and health?
- Is the positive action of ZnO on piglet gut health only due to antibacterial activity?
- Do all sources of zinc oxide perform equally?

## Mechanisms of ZnO

Zinc oxide is a non-water soluble compound which can dissociate at low pH. The gastric environment of young pigs is not acidic enough for full dissociation with immature digestive system and too high buffer capacity of the feed.

Once dissociated, solubilised Zn<sup>2+</sup> are released into the gastro-intestinal tract. It is generally thought that the antimicrobial properties of ZnO are only attributed to these ions, which interact with bacterial amino acids and play a role in the inhibition of active transporters.

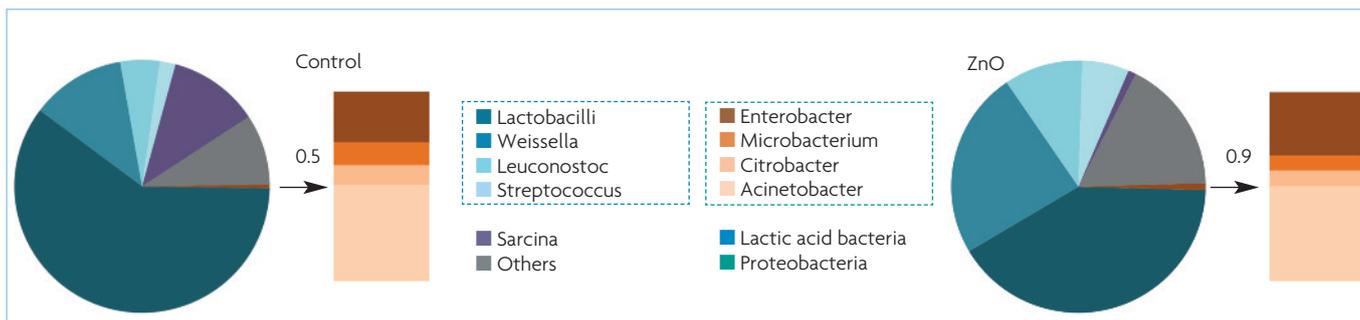
However, if the antibacterial activity was limited to Zn<sup>2+</sup>, water soluble zinc sources like zinc sulphate would be more efficient.

Recent studies have confirmed a specific effect from ZnO molecule. The contribution of soluble zinc ions to the antimicrobial activity of ZnO differed among bacterial species and can be very low: for example, only 15% for Escherichia coli (Fig. 2).

This would explain why piglet diets are only supplemented with pharmacological dosage of zinc oxide and not with other zinc compounds. It is still debated if the general positive effect of ZnO on piglet gut health can only be attributed to antibacterial activity. Among the most likely hypotheses concerning the specific effect of ZnO, this zinc source would be able to generate reactive oxygen species (ROS), under UV exposure but also in dark-

*Continued on page 13*

**Fig. 1. Effect of the pharmacological dosage of ZnO on the bacteria distribution in the intestine.**



Continued from page 11  
and consequently causes severe damage to the cells.

ROS induce oxidative stress and can cause the oxidation of bacterial lipid membrane; this could lead to an alteration of the membrane composition and partially explain the reduced adherence of the pathogenic bacteria like *E. coli* to host cells.

This reduction of the bacteria attachment to the epithelium would avoid the secretion of enterotoxins, leading to diarrhoea.

Variable effects have been observed from ZnO supplementation on gut morphology.

Currently, mechanisms of the antimicrobial activity of ZnO are not totally identified and could vary with the target cells, but it is recognised that specific mechanisms exist. The mechanisms by which zinc oxide products will improve piglet intestinal health can also depend on the chemical features of the particles in the powder.

### Not all zinc oxides are equal

There are many different zinc oxide sources. Any result obtained with analytical grade trace minerals, which are standardised, can not be fully extrapolated to feed grade sources, which are highly variable. Zinc oxide can be manufactured with various raw materials and manufacturing processes.

Physico-chemical properties of ZnO products used in animal nutrition show strong differences, which explain variable bioavailability values and antibacterial strength.

Manufacturing zinc oxide at the nano size can elicit significant differences compared to the bulk equivalent and may be of potential interest in livestock feeding.

However, before testing engineered nano ZnO particles in animal diets, such challenges will need to be addressed: product characterisation with appropriate analytical methods, industrial scaling from lab-



Various feed grade ZnO sources and potentiated zinc oxide (centre).

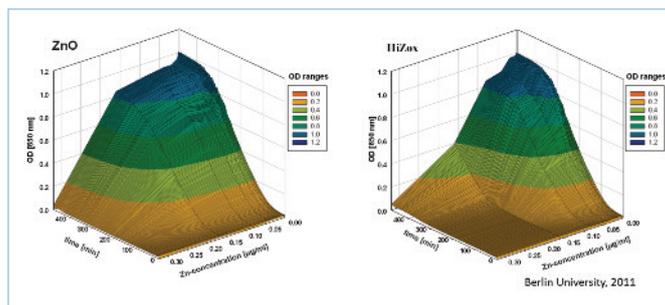


Fig. 3. Effects of regular zinc oxide and HiZox on the growth kinetics of pathogenic strains of *E. coli* in neutral condition.

oratory samples, ADME (Absorption, Distribution, Metabolism, Excretion) and toxicity studies on animals, safety for workers and for the environment, and compliance with local regulation.

### Potentiated zinc oxide

A patented manufacturing technology is at the origin of a potentiated form of zinc oxide, HiZox, which has been developed for animal nutrition.

Physico-chemical properties are modified: specific particle size and shape with increased (10-15 times higher than conventional sources) specific surface area drastically

increase the surface of contact with bacteria. The high porosity of the powder amplifies the antibacterial activity of this potentiated zinc oxide. Scientific experiments have confirmed its efficacy to manipulate intestinal and faecal microbiota of weaned piglets.

### In vitro bacterial growth

Studies supervised by the University of Berlin have compared the inhibitory effects of HiZox and regular zinc oxide on the growth of pathogenic strains of *Escherichia coli* (fimbriated *E. coli* PS79/K88/F4 and PS7/K81/F18). HiZox showed stronger inhibitory action on *E. coli* in neutral (pH 6.5) and acidic conditions (pH 4.6) which mimic gastrointestinal pH environment (Figs. 3, 4).

Fig. 4. Effects of regular ZnO and HiZox on the lag times of pathogenic strains of *E. coli* in acidic conditions.

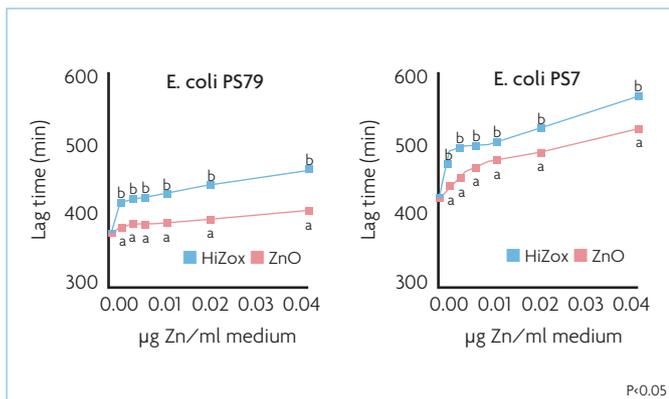
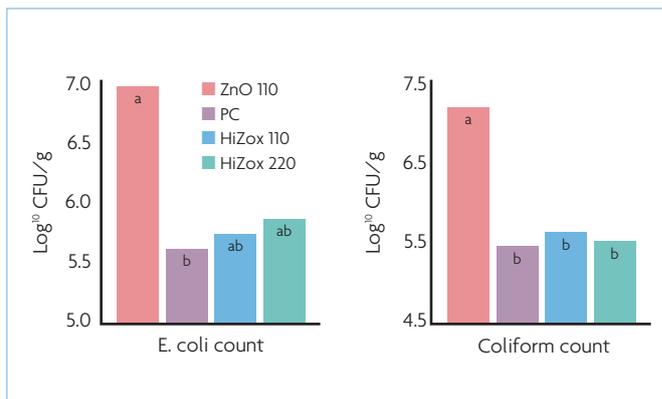


Fig. 5. *E. coli* count (left) and coliform count (right) in the distal small intestine.



### Intestinal microbial populations

A study performed in Ghent University (Michiels, 2016) measured bacterial counts of *E. coli* and coliforms in the small intestine of weaned piglets, which were fed one of the four experimental diets: negative control (110mg/kg Zn from standard ZnO); positive control (3kg/T ZnO); HiZox at EU legal dosage (110mg/kg Zn) and higher dosage (220mg/kg Zn). Replacing the conventional zinc oxide by HiZox decreased microbial concentrations in the small intestine (Fig. 5).

### Ex vivo bacterial growth

Optimal manipulation of gut microbiota in piglets is not only reducing pathogenic bacteria, but also decreasing bacterial load in the small intestine. An experiment (Vahjen et al, JAS 2012) compared the growth repressing effect of HiZox and regular zinc oxide on bacteria from stomach and jejunum samples from weaned piglets. Differences in the small intestine (Fig. 6) are likely to originate in the stomach (Fig. 7).

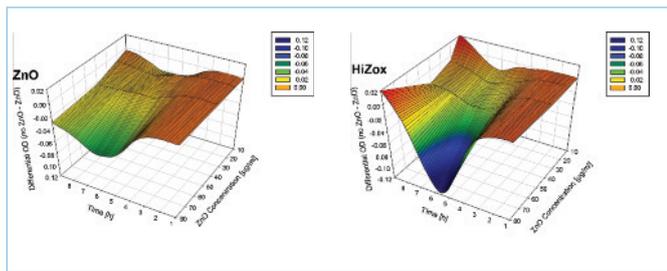
### Faecal E. coli pathogenic factors

An experiment conducted by Berlin University (Vahjen, JAS 2016) analysed via PCR assays total enterobacteria, the *Escherichia* group and *E. coli* pathogenic factors in faecal samples from weaned piglets which were fed one of the four experimental diets: negative control (110mg/kg Zn from standard ZnO); positive control (3kg/T ZnO); HiZox at EU legal dosage (110mg/kg Zn) and higher dosage (220mg/kg Zn). More than half of the samples in the negative control were positive for at least one of the analysed *E. coli* pathogenic factors, but the other treatment groups showed much lower percentages of positive samples (Fig. 8).

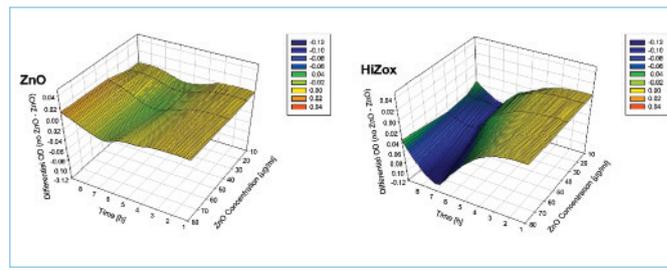
### Faecal microbial populations

One study supervised by Dankook University (Cho et al, Anim. Sci. Journ. 2015) measured the effect of zinc oxide dose and source on growth performance, nutrient digestibility, blood profiles, faecal

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**Fig. 6. Bacterial growth inhibition of ZnO and HiZox, in the stomach.**



**Fig. 7. Bacterial growth inhibition of ZnO and HiZox, in the jejunum.**

Continued from page 13  
microbial shedding and faecal score of weaned piglets which were fed one of the four experimental diets: negative control (110mg/kg Zn from the premix); positive control (3kg/T ZnO); HiZox at two levels (200mg/kg and 300mg/kg).

It was concluded that HiZox modified the faecal Lactobacilli/*E. coli* ratio (Fig. 9), thus improving faecal score and growth performance.

### Conclusion

There is growing pressure on the reduction of antibiotics and pharmacological dosage of zinc oxide in piglet diets. Excessive and prolonged used of ZnO are known for negative consequences: risk of heavy metal contaminations, nutri-

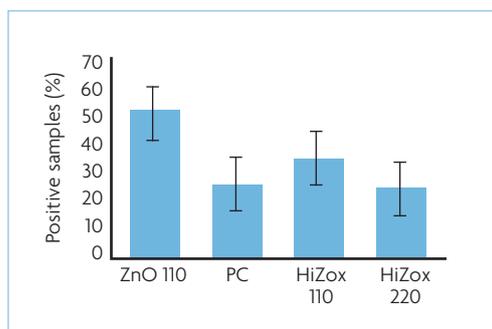
tional antagonisms, environmental accumulation, bacterial resistance. Feed formulators are seeking feed ingredients and feed additives which can reduce the risk of diges-

tive disorders in critical phases like the post-weaning period. HiZox can be used, in combination with alternatives to AGPs, in a preventive way to improve intestinal health and

reduce the need for non-sustainable practices. ■

References are available from the author on request

**Fig. 8. Percentage of faecal samples positive for *E. coli* pathogenic factor.**



**Fig. 9. Ratio of faecal lactobacilli/*E. coli* at 14 days post weaning.**

