EFFECT OF DIETARY SUPPLEMENTATION WITH α-TOCOPHERYL ACETATE ON THE STABILITY OF LOW NITRITE COOKED TURKEY HAM.

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Background Lipid oxidation is one of the main causes of deterioration in the quality of meat and meat products during storage (Tichivangana & Morrissey., 1985). The process occurs as a result of the reaction between atmospheric oxygen and unsaturated fatty acids. Peroxides are intermediate products in the oxidation process, which in turn break down to odour and flavour producing compounds such as aldehydes, ketones, and allegated the products of the process of th alcohols etc. (Rendl et al., 1982). The flavour and odour of this mixture renders the meat unacceptable for human consumption.

Nitrite has been found to be effective in preventing lipid oxidation and is also responsible for desirable colour formation in cured meat products (Rubin., 1977). However in recent years there has been some controversy over the use of nitrite due to nitrosamine formation. During heating residual nitrite may react with certain amines to produce nitrosamines. These compounds are considered to be carcinogenic (Sen et al., 1973; Gray et al., 1982). The maximum level of nitrite allowed in meat is 200 ppm in Ireland and 120 ppm in the U.S.A (Tichivangana et al., 1984). Therefore, it is desirable to reduce the nitrite levels in meat. α-Tocopherol has shown good antioxidant properties (Buckley & Connolly.,

1980), and therefore could be used to replace some of the nitrite in processed meats.

The objective of this study was to investigate the effect of reducing added nitrite levels in cooked turkey ham produced from meat containing high and low levels of dietary α-tocopheryl acetate.

Experimental Design

In this study, turkeys were fed diets containing either low α-tocopheryl acetate of 20 mg/kg or high supplemented α-tocopheryl acetate (600 mg/kg) from 1 week old up to slaughter at 21 weeks. The breast meat was manufactured into cooked turkey ham using commercial manufacturing procedures. The input levels of nitrite in the cooked turkey ham were 0 ppm, 50 ppm, and 100 ppm. The effect of storage in a display cabinet, illuminated under fluorescent light (616 LUX) at 4°C on vacuum packed (30-100 cm³/m²/24 hours) and over-wrapped oxygen Permeable film (6000-8000 cm³/m²/24 hours) samples was determined by measuring oxidative (TBARS) and colour (Hunter "a" values) changes.

Materials and Methods

α-TOCOPHEROL ANALYSIS

Performed by High performance liquid chromatography. The extraction procedure is based on the method of Butriss and Diplock (1984), while, the HPLC method is a modification of Bieri et al. (1979).

THIOBARBITURIC ACID (TBARS)

Performed according to the method of Tarladgis et al. (1960)
COLOUR ANALYSIS

Analysed by the Hunter L, a, b system, using the minolta colorimeter (Model: -CR 300)

The residual α -tocopherol level in the raw turkey breast meat from birds fed the supplemented diet was 5 μ g/g by comparison to 1 μ g/g for breast meat from the basal group. In the oxygen permeable overwrapped samples (FIG. 1 & 2) and the vacuum packaged product (FIG. 3 & 4) the α -tocopherol level in the raw turkey breast meat from the basal group. In the oxygen permeable overwrapped samples (FIG. 1 & 2) and the vacuum packaged product (FIG. 3 & 4) the α -tocopherol level in the raw turkey breast meat from the basal group. In the oxygen permeable overwrapped samples (FIG. 1 & 2) and the vacuum packaged product (FIG. 3 & 4) stream the same general trends were detected. However, the differences in the vacuum packaged product were not as evident. At low levels of α-locopherol the product with 50 ppm NO₂ had poor colour stability by comparison to the product with 100 ppm NO₂ (FIG. 2). The product with high α-tocopherol and 50 ppm input NO₂ had improved colour stability over time compared to the product with 50 ppm NO₂ at low levels of α-tocopherol. At low levels of α-tocopherol the oxidative stability was lower in the product with 50 ppm NO₂ than in the product with 100 ppm input NO₂ (FIG. 2). Ppm input NO₂ (FIG. 1). It was found that at high α -tocopherol levels the oxidative stability of the product with 50 ppm added NO₂ showed no difference to the product with 100 ppm input NO₂ at the same level of α -tocopherol up to day 6 (FIG. 1). At high α -tocopherol levels the product with 50 ppm input NO₂ showed much better oxidative stability over time than the product with the 50 ppm nitrite at low α - tocopherol levels.

Conclusions

At high α-tocopherol levels the over-wrapped cooked turkey ham with 50 ppm input NO₂ had better oxidative stability and better colour stability than the product with 50 ppm NO₂ at low α-tocopherol levels

The way local and 50 ppm input NO₂ had better oxidative stability and colour stability

The vacuum-packed cooked turkey ham containing high α -tocopherol and 50 ppm input NO₂ had better oxidative stability and colour stability than the product with low a-tocopherol and 50 ppm input NO₂.

Reduction of input nitrite levels from 100 ppm to 50 ppm at high α-tocopherol levels maintains colour and oxidative stability for a longer period of time.

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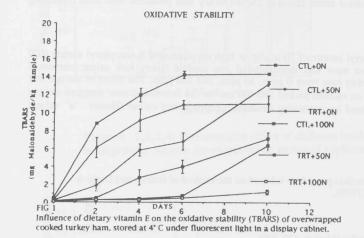
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6 VALUES TRT+50N 4 0 HUNTER TRT+100N - 2 TRT+ON CTL+100N CTL+ON - 4 CTL+50N DAYS 4 FIG 2 0 Influence of dletary vitamin E on the colour stability of over-wrapped cooked turkey ham, stored at 4° C under fluorescent light in a display cabinet

COLOUR STABILITY

