Atheromatous Changes in Cholesterol-Fed Rabbits Treated with Eggplant Extract


Cholesterol-induced changes in rabbits were studied with and without administration of an extract of eggplant. An anemia, accompanied by atrophic gastritis, developed in the cholesterol-fed animals. Their atheromatous plaques, and elevated serum cholesterol, lipoprotein, a-lipoprotein, and neutral fat were not affected by the eggplant extract.

It was reported by Roffo 1 that feeding dried eggplant (Solanum melongena L.), or injecting an alcoholic extract of the dried vegetable subeutaneously, caused a fall in the blood cholesterol levels of rabbits. The variation in the analytic values reported rendered the interpretation somewhat questionable. Later Herrmann 2 found that eggplant, although less effective than choline, methionine, or inositol, lowered the blood cholesterol levels when administered to hens or to hypercholesteremic patients. Roffo 1 attributed this action of eggplant to its cholagogue action. Later both Noble 3 and Trelles Duelo 4 confirmed the claim that eggplant is a cholagogue.

On the other hand, Wilkinson, Jackson, and Vogel 5 in a test on 6 healthy male subjects, were unable to demonstrate any effect on serum cholesterol levels from feeding the dried equivalent of 180 to 360 gm. of fresh eggplant daily. In view of these conflicting findings and of the present interest in hypercholesteremia and its control, it was decided to reinvestigate this decholesterolizing action of eggplant.

Methods

Forty rabbits (38 males and 2 females), varying in weight from 1065 to 3710 Gm., and of mixed breeding, were listed in order of increasing weight, and subdivided into successive lots of five. With the aid of sets of random numbers, one rabbit was assigned from each of these lots to each of five treatment groups. In this manner the initial mean weights of the treatment groups were made approximately equal. The rabbits were housed in individual cages in an air-conditioned room. Wood shavings were used on the floors of the cages.

During the one week pre-experimental period, water and Purina rabbit feed in pellet form were allowed ad libitum. For the last three days the pelleted feed was removed in the evening. In the morning, the rabbits were given approximately 25 Gm. of ground carrots which they had to consume before the pelleted feed was returned.

At the end of the pre-experimental period, blood samples of about 3 ml. were taken from the ear vein of each rabbit after 16 hours' fasting. The rabbits were allowed feed until evening and again starved overnight. On the following morning they were given approximately 25 Gm. of ground carrots containing either (a) no addition (diet I), (b) 1 Gm. of cholesterol U.S.P. (diet II), (c) approximately 6 Gm. of eggplant extract (diet III), (d) 1 Gm. of cholesterol plus 6 Gm. of eggplant extract (diet IV), or (e) as (b) for the first 5 weeks and as (d) for the last 4 weeks. None of the rabbits was fed regular pelleted diet until all had eaten the supplement, in which it was occasionally necessary to mix a few ground pellets to improve palatability. The supplement usually was consumed in 2 to 3 hours, after which all animals received pelleted feed until it was again removed in the evening. This regimen was followed 7 days a week except on every second Monday, when blood samples were taken and body weights recorded.

Blood samples were held at room temperature for one hour, refrigerated for one hour, and centrifuged. A portion of the serum was stored in the frozen state in capped tubes for subsequent cholesterol determinations, and the remainder was stored at 4 C. for use in electrophoretic studies within the week.

Preparation of Eggplant Extract. Fresh eggplants were trimmed, chopped into conveniently sized pieces, and in 2 Kg. lots, blended with 1 L. of ethanol in a large Waring Blender until thoroughly macerated. The slurry was strained...
through several layers of cheesecloth and the residue was squeezed as dry as possible. The residues from two such extractions were combined and reextracted in the blender with a second liter of alcohol. Again the residue was removed with the aid of cheesecloth. The combined filtrates were allowed to stand overnight and then filtered through two layers of Whatman No. 1 filter paper. Some 500 L. of extract were obtained from 245 Kg. of fresh, trimmed eggplant. This extract was concentrated in vacuo in a Flash Evaporator with continuous addition of extract and a bath temperature of 55 to 60 C. The final, thick, brown syrup occupied 9,700 ml. and had a specific gravity of 1.36. The finished concentrate was thoroughly mixed and stored in the refrigerator until required. The material was conveniently dispensed with the aid of a large syringe, and 4.5 ml., corresponding to approximately 6 Gm., were mixed with the supplemental feeding of ground carrot per rabbit as required. This amount of extract was approximately equivalent to 110 Gm. of fresh eggplant.

Electrophoretic Separation of Serum Proteins and Lipoproteins. Serum samples were subjected to electrophoresis on agar with a slight modification of the apparatus and procedure by Giri.6 A diagram of the apparatus is shown in figure 1. The agar was prepared as a 1 per cent solution in pH 8.6 veronal acetate buffer of ionic strength 0.05. The sample of serum, 10L, or 20A, was pipetted onto a strip of 3 MM Whatman paper 20 mm. long and 1 or 2 mm. wide and placed on the agar film. A potential of 150 volts, equivalent to approximately 4 volts/cm., was applied to the films, giving a current flow of approximately 3.5 ma. per film at room temperature. After three hours the current flow was cut off, the cover plates were removed, and the films were dried in a current of air. The filter paper connector strips were removed, and the cellophane bearing the dried agar film was stripped from the frame, suitably identified, and stored in a dry place until stained.

Staining. (a) Protein: Anido black was purified according to the method of Kawerau.7 The dye solution was prepared by dissolving 5 Gm. of dye in a mixture of 600 ml. of distilled water, 400 ml. of methanol, and 100 ml. of glacial acetic acid. The strips were immersed in the dye solution for 30 minutes and were then washed in three to four changes of 10 per cent acetic acid in methanol until the background was clear. The strips were pressed between dry filter papers. (b) Lipoprotein: The films were immersed overnight in a saturated solution of Oil Red O in 50 per cent ethanol. The stained strips were washed in three or four changes of 50 per cent ethanol and pressed between dry filter papers. According to Uriel and Grabar8 and the present findings, unlike the re-
sults of paper electrophoresis, no lipoprotein with a mobility similar to that of the B-globulins is found. Rather, most of the mobile lipid is associated with the albumin and a-protein fractions.

**Interpretation.** The 10% sample patterns, stained for protein, were scanned in the densitometer with light of 615 mm wavelength. The optical densities were plotted against distance in mm. on graph paper. The areas enclosed by the curves were cut out with scissors, subdivided as indicated into one or more fractions, and weighed for a roughly quantitative relative measure.

**Cholesterol.** Serum cholesterol was determined by the Henly modification of the method of Zlatkis, Zak, and Boyle.

Cholesterol determinations were made on the left adrenal of each rabbit. The weighed adrenal was ground in a Potter and Elvehjem type of homogenizer in a small volume of the Henly reagent (0.05 per cent Fe Cl₃. 6H₂O in glacial acetic acid). The homogenized tissue suspension was transferred to a graduated centrifuge tube and made to volume with the reagent. After centrifuging and filtering through glass wool, a suitable aliquot of the filtrate was transferred to a glass-stoppered centrifuge tube.

**Liver Lipid.** The procedure used for the determination of liver lipid was that of Folch et al. Because of the alternatives permitted, the specific method is presented in detail. A 2.5 Gm. sample of liver was homogenized in CHCl₃:MeOH 2:1, transferred to a 50 ml volumetric flask, and the flask was filled to the mark with solvent. After mixing, the flask contents were filtered through No. 42 Whatman paper and 20 ml aliquots were transferred to glass-stoppered centrifuge tubes. After the addition of 4 ml of 0.04 per cent aqueous CaCl₂ solution and thorough mixing, the tubes were centrifuged. The upper layer was removed as completely as possible with the aid of suction, and the remaining lower phase was washed three times with a mixture of CHCl₃: MeOH:H₂O in the proportions 3:48:47. The lower phase was then treated with a little methanol to make a single phase, and poured into tared flasks. The washings, made with CHCl₃:MeOH: H₂O in the proportions 86:14:1, were added, and the whole was evaporated to dryness in the water bath in vacuo at 60 C. The dried flasks were further dried in desiccators in vacuo and weighed.

**Histologic Technics.** Tissues taken for examination were fixed in 10 per cent formalin in normal saline. Sections were cut at 4μ and were stained with hematoxylin and eosin. Sections of aorta were also stained with Verhoeff's elastica stain. Lillie's modification of the ferrocyanide reaction of M. Perls was used on kidney and spleen sections.

**Hematologic Technics.** Red blood cell counts were made on a Spencer Bright-line hematocytometer. The Van Allen hematocrit method was used to determine the packed cell volume. Hemoglobin estimations were made with a slight modification of the pyridine-hemochromogen method of Rimington. Blood and bone marrow smears were stained with Giemsa.

After the animals had been 12 weeks on experiment, the final blood samples were taken, and the red cell count, the hematocrit and hemoglobin values were determined, and a blood smear was made. The rabbits were then decapitated and examined for gross pathology. A bone marrow smear was prepared, the liver and left adrenal were weighed, and portions of the liver and the adrenal were placed in sealed vials and frozen. Samples of liver, spleen, stomach, kidney, prostate, testes, pancreas, right adrenal, and heart with attached major vessels, were preserved for examination.

**RESULTS**

**Growth and Mortality.** In table 1 are recorded the mean body weights of the five groups of rabbits throughout the test. The high standard deviations reflect the range of

<table>
<thead>
<tr>
<th>Weeks on diet</th>
<th>Diet</th>
<th>1</th>
<th>2</th>
<th>3</th>
<th>4</th>
<th>5</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>I</td>
<td>2119±779</td>
<td>2085±611</td>
<td>1992±571</td>
<td>2041±659</td>
<td>2140±694</td>
</tr>
<tr>
<td>2</td>
<td>II</td>
<td>2355±689</td>
<td>2266±525</td>
<td>2223±487</td>
<td>2246±569</td>
<td>2337±651</td>
</tr>
<tr>
<td>4</td>
<td>III</td>
<td>2453±658</td>
<td>2414±417</td>
<td>2411±405</td>
<td>2448±506</td>
<td>2501±549</td>
</tr>
<tr>
<td>6</td>
<td>IV</td>
<td>2588±605</td>
<td>2488±362</td>
<td>2406±355</td>
<td>2564±459</td>
<td>2563±536</td>
</tr>
<tr>
<td>8</td>
<td>V</td>
<td>2733±562</td>
<td>2588±339</td>
<td>2651±340</td>
<td>2660±486</td>
<td>2691±543</td>
</tr>
<tr>
<td>10</td>
<td>I (Survivors)</td>
<td>2808±540</td>
<td>2584±385</td>
<td>2686±340</td>
<td>2766±483</td>
<td>2641±562</td>
</tr>
<tr>
<td>12</td>
<td>II</td>
<td>2901±505</td>
<td>2658±378</td>
<td>2786±385</td>
<td>2800±509</td>
<td>2724±635</td>
</tr>
</tbody>
</table>

*Survivors.*
body weights in the animals at the start of
the experiment, an unavoidable circumstance
because of the difficulty of procuring rabbits.
It is easily seen, however, that the treatments
had no effect on the body weight of the test
animals. Survival figures also are noted in
this table. One animal on each of diets II,
III and IV died during the experiment, under
circumstances that in no instance could be di-
rectly related to the diets. Two succumbed
to acute Pasturella infection. The cause of
death in the third animal was not deter-
dined.

Serum Protein and Lipoprotein. The elec-
троphoretic pattern suggested that feeding
cholesterol tended to increase the β-globulin
and decrease the γ-globulin fractions of rabbit
blood serum. The individual variations were
such that these differences did not prove to be
statistically significant. The feeding of egg-
plant extract did not appear to have any in-
fluence on the results. It might be mentioned
here that the technic used (electrophoresis in
agar) leads to a higher percentage of protein
designated as albumin than is the case when
paper is used as the supporting medium. In
part, at least, this may be the result of con-
siderable "trailing" of albumin on paper and
the lack of this effect in agar.

In the case of lipids, feeding cholesterol
caused a tremendous increase in material
stainable by Oil Red O, an increase which be-
came progressively greater as the experiment
continued. The changes occurred in the lipo-
albumin, in the α-lipoprotein, and in the neu-
tral fat. The data, for total stainable material,
in arbitrary units, are presented in table 2.
It is apparent that feeding of eggplant extract
along with the cholesterol or during the last
third of the experimental period had no sig-
nificant influence on the lipid level attained.

Serum Cholesterol. Feeding cholesterol
caued a prompt increase in serum cholesterol
levels which tended to show a further gradual
increase as feeding was prolonged. The pre-
experimental serum cholesterol levels were
quite variable, and this variability persisted
when the animals were fed cholesterol. The
administration of eggplant extract appeared
to have no effect on serum cholesterol levels
(table 3).

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**Table 2.—Oil Red O Stainable Material in Serum Electropherograms; Arbitrary Units ±
Standard Deviation**

<table>
<thead>
<tr>
<th>Weeks on diet</th>
<th>I</th>
<th>II</th>
<th>III</th>
<th>IV</th>
<th>V</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>244±74</td>
<td>265±31</td>
<td>319±124</td>
<td>269±99</td>
<td>196±59</td>
</tr>
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<td>2</td>
<td>227±53</td>
<td>446±133</td>
<td>284±90</td>
<td>373±25</td>
<td>389±129</td>
</tr>
<tr>
<td>4</td>
<td>201±44</td>
<td>548±140</td>
<td>276±74</td>
<td>514±168</td>
<td>565±140</td>
</tr>
<tr>
<td>6</td>
<td>288±62</td>
<td>631±151</td>
<td>289±82</td>
<td>638±152</td>
<td>550±108</td>
</tr>
<tr>
<td>8</td>
<td>279±56</td>
<td>711±200</td>
<td>344±173</td>
<td>631±193</td>
<td>611±141</td>
</tr>
<tr>
<td>10</td>
<td>265±49</td>
<td>745±160</td>
<td>315±127</td>
<td>776±127</td>
<td>642±154</td>
</tr>
<tr>
<td>12</td>
<td>302±91</td>
<td>789±249</td>
<td>364±166</td>
<td>862±234</td>
<td>672±223</td>
</tr>
</tbody>
</table>

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**Table 3.—Serum Cholesterol Mg./100 Ml.**

<table>
<thead>
<tr>
<th>Weeks on diet</th>
<th>I</th>
<th>II</th>
<th>III</th>
<th>IV</th>
<th>V</th>
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<tbody>
<tr>
<td>0</td>
<td>70±46</td>
<td>99±63</td>
<td>73±25</td>
<td>49±27</td>
<td>96±84</td>
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<tr>
<td>2</td>
<td>77±56</td>
<td>646±357</td>
<td>84±51</td>
<td>557±396</td>
<td>714±546</td>
</tr>
<tr>
<td>4</td>
<td>76±39</td>
<td>684±496</td>
<td>67±26</td>
<td>607±390</td>
<td>752±325</td>
</tr>
<tr>
<td>6</td>
<td>50±21</td>
<td>1105±539</td>
<td>96±99</td>
<td>967±520</td>
<td>1228±395</td>
</tr>
<tr>
<td>8</td>
<td>32±15</td>
<td>1730±423</td>
<td>57±31</td>
<td>1340±785</td>
<td>1773±758</td>
</tr>
<tr>
<td>10</td>
<td>48±18</td>
<td>1779±617</td>
<td>80±55</td>
<td>1504±793</td>
<td>1831±749</td>
</tr>
<tr>
<td>12</td>
<td>31±10</td>
<td>1696±548</td>
<td>38±17</td>
<td>1457±855</td>
<td>1764±784</td>
</tr>
</tbody>
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**ATHEROMATOUS CHANGES AND EGGPLANT EXTRACT**

**Table 4.** Adrenal Cholesterol

<table>
<thead>
<tr>
<th>Diet</th>
<th>Adrenal weight (mg.)</th>
<th>Cholesterol (mg./adrenal)</th>
<th>% Adrenal</th>
</tr>
</thead>
<tbody>
<tr>
<td>I</td>
<td>160± 57</td>
<td>11.7± 7.5</td>
<td>7.26± 2.9</td>
</tr>
<tr>
<td>II</td>
<td>340± 77</td>
<td>25.4± 3.5</td>
<td>7.53± 2.2</td>
</tr>
<tr>
<td>III</td>
<td>189± 65</td>
<td>16.0± 6.2</td>
<td>7.99± 2.8</td>
</tr>
<tr>
<td>IV</td>
<td>39± 75</td>
<td>24.4± 3.1</td>
<td>6.66± 2.0</td>
</tr>
<tr>
<td>V</td>
<td>339±115</td>
<td>25.4± 1.8</td>
<td>6.97± 2.3</td>
</tr>
</tbody>
</table>

**Adrenal Cholesterol.** Adrenal total cholesterol was significantly greater in amount in rabbits fed cholesterol (table 4). Feeding of eggplant extract had no influence on the results whether alone or with cholesterol. When the adrenal cholesterol content was calculated on a percentage basis it was apparent that the concentration in the adrenals was not significantly different between diets. The apparent differences in total adrenal cholesterol was caused by the hypertrophy of the adrenals of cholesterol-fed rabbits.

**Liver Lipid.** The alterations in liver weights caused by feeding cholesterol are shown, together with the lipid content in table 5. The increases in liver weight on diets II, IV, and V were significant only at \( p = 0.1 \). At the same time, this enlargement, when considered in conjunction with the very significant elevation in liver lipid concentration, indicated that feeding cholesterol resulted in a very marked increase in total liver lipid. Feeding eggplant extract appeared to have no influence on liver weight or lipid concentration.

**Blood Picture.** The results of the various determinations are presented in table 6. It is apparent that feeding cholesterol to rabbits caused a significant decrease in the red cell count, in hemoglobin concentration, and in percentage of packed cells. The mean corpuscular volume was not significantly altered, although there was some suggestion that the value was elevated in the cholesterol-fed animals. The mean corpuscular hemoglobin concentration was not affected. The color index was normal. The blood picture in the cholesterol-fed rabbits showed anisocytosis, poikilocytosis and polychromatophilia, megaloblasts, and normoblasts. “Tailed” and “racquet” erythrocytes were also noted.

In no instance did there appear to have been any influence on the blood picture from feeding eggplant extract. These results led to the conclusion that feeding cholesterol caused a macrocytic anemia in rabbits.

**Gross Pathology**

**Atheroma Index.** The feeding of cholesterol to rabbits had a profound effect on the incidence and severity of plaque formation in the rabbit aorta. Eggplant extract had no effect.

**Table 5.** Liver Lipid

<table>
<thead>
<tr>
<th>Diet</th>
<th>Liver wt. (Gm.)</th>
<th>% Lipid</th>
</tr>
</thead>
<tbody>
<tr>
<td>I</td>
<td>119±25</td>
<td>4.04±0.45</td>
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<tr>
<td>II</td>
<td>158±32</td>
<td>8.75±1.16</td>
</tr>
<tr>
<td>III</td>
<td>120±21</td>
<td>4.53±0.78</td>
</tr>
<tr>
<td>IV</td>
<td>155±28</td>
<td>9.81±2.81</td>
</tr>
<tr>
<td>V</td>
<td>154±56</td>
<td>8.15±0.64</td>
</tr>
</tbody>
</table>

**Table 6.** Hematology

<table>
<thead>
<tr>
<th>Diet</th>
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<th>II</th>
<th>III</th>
<th>IV</th>
<th>V</th>
</tr>
</thead>
<tbody>
<tr>
<td>R.B.C. millions/mm$^3$</td>
<td>6.7±0.6</td>
<td>4.4±1.0</td>
<td>6.1±0.8</td>
<td>4.2±1.0</td>
<td>4.0±1.1</td>
</tr>
<tr>
<td>Hemoglobin Gm./100 ml.</td>
<td>14.5±0.7</td>
<td>12.8±2.5</td>
<td>14.5±1.0</td>
<td>11.1±2.8</td>
<td>10.0±2.9</td>
</tr>
<tr>
<td>Hematocrit % packed cells</td>
<td>40.1±5.6</td>
<td>32.5±6.9</td>
<td>41.3±5.0</td>
<td>35.7±7.2</td>
<td>28.6±5.6</td>
</tr>
<tr>
<td>M.C.V.</td>
<td>60.2±8.2</td>
<td>72.6±10.9</td>
<td>67.9±8.3</td>
<td>81.3±12.3</td>
<td>75.6±23.3</td>
</tr>
<tr>
<td>M.C.H.C.</td>
<td>36.7±7.0</td>
<td>37.7±1.7</td>
<td>33.7±2.7</td>
<td>32.8±2.6</td>
<td>35.0±6.3</td>
</tr>
<tr>
<td>Color index</td>
<td>0.72</td>
<td>0.93</td>
<td>0.78</td>
<td>0.88</td>
<td>0.82</td>
</tr>
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</table>
Table 7.—Atheroma Indices

<table>
<thead>
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<th>Diet</th>
<th>Index</th>
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</thead>
<tbody>
<tr>
<td>I</td>
<td>0</td>
</tr>
<tr>
<td>II</td>
<td>12.4 ± 8.8</td>
</tr>
<tr>
<td>III</td>
<td>0</td>
</tr>
<tr>
<td>IV</td>
<td>10.6 ± 7.8</td>
</tr>
<tr>
<td>V</td>
<td>12.8 ± 8.0</td>
</tr>
</tbody>
</table>

and Tejada,15 The aorta was divided into two areas for evaluation. Area 1 was 3 cm. long and included the aortic arch and a small portion of descending aorta. Area 2 was the remainder of the aorta to the aortic bifurcation. The aorta was opened on the dorsal surface, pinned flat on a block of wood with the intimal surface up, and was examined by the naked eye and with a stereoscopic microscope. Three grades only were used as follows: grade I, lipid patches, grade II, lipid streaks, and grade III, fibrous and lipid plaques. (Grouping was made according to the method of Gore and Tejada.) The indices so derived are presented in table 7.

Coccidial abscesses occurred in the livers of 4 control and 6 test animals.

Parasitic cysts (Cysticercus pisiformis) were present in the abdominal cavity of 2 test animals.

Histopathology. Microscopic changes in the aortas of the animals receiving cholesterol and eggplant extract were similar.

Clusters of foam cells were noted in the kidney and spleens of all except the control and eggplant extract group.

Several animals on diets II, IV, and V exhibited an atrophic gastritis. Arteries in the mucosa and submucosa of these animals were partially occluded with foam cells. The atrophy was most obvious in the pyloric portion of the stomach. It is interesting to note that this pathology was most marked in the animals that were markedly anemic, and that cholesterol-fed rabbits that were not anemic had no observable gastric atrophy. As the anemia was of the macrocytic type, it may have been related to the atrophic gastritis, although mucosal cells of the fundus were not obviously abnormal. Studies are being conducted to assess the validity of this theory.

A squamous cell carcinoma was observed in the stomach of 1 animal on diet IV.

The liver and adrenal changes were similar to those described by other workers.

Summary

Feeding cholesterol to rabbits during a 12 week period with or without the addition of eggplant extract had no effect on body weight or mortality and little influence on the serum protein level or distribution. Rabbits fed cholesterol had an elevated serum cholesterol and increased levels of lipoalbumin, α-lipoprotein and neutral fat. They showed an adrenal hypertrophy but no change in the concentration of cholesterol in the adrenals. Liver lipid levels were considerably elevated but liver weights were only slightly increased. Eggplant extract alone or fed with the cholesterol did not alter the results. An anemia developed in rabbits fed cholesterol. The measurements made indicated that this was a macrocytic type of anemia. Gross and microscopic pathological investigations revealed that most of the rabbits fed cholesterol developed atheromatous plaques while incidence in the controls was zero. Eggplant extract appeared not to alter the incidence of plaques.

Cholesterol-fed rabbits that developed anemia were affected with an atrophic gastritis. There appeared to be a relationship between the anemia and gastritis; the more anemic animals exhibited a marked gastric atrophy.

Acknowledgment

The authors wish to thank Dr. W. P. McKinley for his encouragement and continued interest.

Summario in Interlingua

Le administration dietari de cholesterol a conilios durante un periodo de 12 septimanas, con o sin le addition de extracto de melongena, habeva nulle effetto super le pesos corporee e super le mortalitale e pae effetto super le nivello o le distribution del proteinas seral. Conilios recipiente cholesterol dietari habeva elevate nivellos seral de cholesterol e augmentate concentrationes de lipo-albumina, de lipo-
proteina alpha, c de grassia neutre. Iste an­i­ma­les mostravá hypertrophia adrenal sed nulle alteration del nivello de cholesterol in le adrenales. Le nivello de lipido hepatic esseva considerabilmente elevate, sed le pesos del hepate esseva augmentate solmente per leve grados. Extracto de melongena per se o in combination con cholesterol non alterava le resultatos. Le conilios recipiente cholesterol disveloppava anemia. Le mesurationes effectede indicava que il se trattava de un typo macrocytic de anemia. Investigations pathologi­que macro- e microscopic revelava que le majoritate del conilios recipiente cholesterol disveloppava placas atheromatose durante que le incidentia de iste phenomo esseva zero in le animales de controlo. Extracto de melongena non pareva influentiar le incidentia del placas.

Le conilios que disveloppava anemia post le administration de cholesterol esseva afficite per un gastritis atrophic. Il pareva exis­tir un relation inter le anemia e le gastritis. Le plus anemic animales exhibiva plus marcate grados de atrophia gastric.

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Atheromatous Changes in Cholesterol-Fed Rabbits Treated with Eggplant Extract
W. DONALD GRAHAM, JOYCE L. BEARE and HAROLD C. GRICE

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