Phosphorus Supplementation Mitigated Food Intake and Growth of Rats Fed a Low-Protein Diet

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Abstract

Background: Low protein intake is associated with various negative health outcomes at any life stage. When diets do not contain sufficient protein, phosphorus availability is compromised because proteins are the major sources of phosphorus. However, whether mineral phosphorus supplementation mitigates this problem is unknown, to our knowledge.

Objective: Our goal was to determine the impact of dietary phosphorus supplementation on food intake, weight gain, energy efficiency, body composition, blood metabolites, and liver histology in rats fed a low-protein diet for 9 wk.

Methods: Forty-nine 6-wk-old male Sprague-Dawley rats were randomly allocated to 5 groups and consumed 5 isocaloric diets ad libitum that varied only in protein (egg white) and phosphorus concentrations for 9 wk. The control group received a 20% protein diet with 0.3% P (NP-0.3P). The 4 other groups were fed a low-protein (10%) diet with a phosphorus concentration of 0.015%, 0.056%, 0.1%, or 0.3% (LP-0.3P). The rats’ weight, body and liver composition, and plasma biomarkers were then assessed.

Results: The addition of phosphorus to the low-protein diet significantly increased food intake, weight gain, and energy efficiency, which were similar among the groups that received 0.3% P (LP-0.3P and NP-0.3P) regardless of dietary protein content. In addition, phosphorus supplementation of low-protein diets reduced plasma urea nitrogen and increased total body protein content (defatted). Changes in food intake and efficiency, body weight and composition, and plasma urea concentration were highly pronounced at a dietary phosphorus content <0.1%, which may represent a critical threshold.

Conclusions: The addition of phosphorus to low-protein diets improved growth measures in rats, mainly as a result of enhanced energy efficiency. A dietary phosphorus concentration of 0.3% mitigated detrimental effects of low-protein diets on growth parameters. Curr Dev Nutr 2017;1:e000943.

Introduction

Protein deficiency can lead to various adverse health effects at any life stage. Protein restriction compromises growth and muscle maintenance in adolescents and leads to weight loss, reduced subcutaneous fat, increased susceptibility to infection, general lethargy, and delayed wound healing in adults (1). In addition, inadequate protein content in maternal diets during pregnancy is reported to result in negative health outcomes in offspring (2). Experiments in rodents have shown that a maternal low-protein diet during pregnancy is associated with disproportionate patterns of fetal growth (3), increased adult onset of metabolic disorders, and increased adiposity and glucose intolerance (4). Furthermore, low dietary protein intake has consistently been associated with increased body fat content in rodents (5–8).

Alterations in food intake (9, 10), organ size, and enzymatic function (11, 12) are believed to mediate animals’ metabolic response to protein restriction. Changes in food

Keywords: low-protein diet, phosphorus, weight gain, food intake, energy efficiency, nonalcoholic fatty liver disease, Sprague-Dawley rats

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Abbreviations used: H&E, hematoxylin and eosin; LP-0.015P, 10% protein and 0.015% P; LP-0.056P, 10% protein and 0.056% P; LP-0.1P, 10% protein and 0.1% P; LP-0.3P, 10% protein and 0.3% P; NP-0.3P, 20% protein and 0.3% P; ORO, Oil Red O.
intake were reported to be inconsistent; some studies showed an increase (7, 8, 13) and others reported a decrease that was dependent on the extent of protein restriction (10, 14). This inconsistency may imply that other factors are involved in the regulation of food intake.

Although proteins vary in phosphorus content and bioavailability (15), the synergistic relationship between dietary protein and phosphorus intake suggests that dietary phosphorus modulates the effects of protein restriction. Phosphorus is an essential mineral and thus plays an important role in cellular metabolism. It is also integrated within ATP, which acts as a phosphate donor for many metabolic reactions (16). Such reactions are strongly dependent on the body’s phosphorus availability for its production (17, 18), and this is related to dietary phosphorus availability.

Populations whose staple foods are low in protein quantity and quality (e.g., cassava, maize, rice) are most at risk of developing diseases associated with protein malnutrition (e.g., marasmus and kwashiorkor) (19). Low dietary phosphorus results from low protein in the diet, which tends to compound nutritional problems. These populations also consume minimal amounts of animal and dairy products, which are important dietary sources of both high-quality proteins and phosphorus. In contrast, most stable plant-based foods that are also commonly consumed by at-risk populations (e.g., whole wheat or brown rice) have low protein quantity and quality and contain low amounts of bioavailable phosphorus because phosphorus is mainly bound in the form of phytate, which makes it unavailable for absorption (20). Fermentation is known to break down phytate and thus improves the bioavailability of phosphorus (21). Fermentation of yeast for ~60 min was reported to reduce the phytate content of bread by ~10% (22).

Phosphorus intake is compromised under conditions of protein restriction; therefore, it can be assumed that inadequate dietary phosphorus intake compounds the negative effects of protein restriction. Accordingly, this study was performed to assess the impact of phosphorus on food intake, weight gain, food efficiency, and body composition of rats maintained with a low-protein diet with varied concentrations of dietary phosphorus.

Methods

Animal housing
Forty-nine 6-wk-old male Sprague-Dawley rats (American University of Beirut Institutional Animal Care and Use Committee). After the 1-wk acclimation period, each rat (weight range: 190–315 g) was randomly allocated to 1 of 5 experimental groups. Under normal conditions, the recommended phosphorus content of the rats’ diet is 0.3% (26), which was the dietary proportion used for the control group. Treatment groups were as follows: 20% protein and 0.3% P (NP-0.3P; control), 10% protein and 0.015% P (LP-0.015P), 10% protein and 0.056% P (LP-0.056P), 10% protein and 0.1% P (LP-0.1P), and 10% protein and 0.3% P (NP-0.3P) (Supplemental Table 1).

The rats were offered their corresponding diets ad libitum for 9 wk. At the study termination, overnight-fastcd rats were anesthetized with isoflurane (Forane; Abbott) and blood was collected from the superior vena cava. The rats were then euthanized by removing the heart and their livers were extracted and weighed. Immediately afterward, 2 small liver sections were taken for histologic analysis and the remaining section was frozen in liquid nitrogen and stored at ~80°C. Blood samples were centrifuged at 2200 × g for 15 min at 3°C and plasma aliquots were collected and stored at ~80°C. The rat carcasses were stored at ~20°C for body composition analysis.

Food intake and body weight and composition
Food intake and body weight were measured twice per week and then averaged to calculate weekly changes. Carcasses were dried until they reached a constant weight (~48 h) at 105°C and homogenized, and fat was extracted using petroleum ether (boiling point: 40–60°C). The moisture and fat content of the carcass was calculated from the differences in weight. Hepatic fat content was determined as follows: ~2 g of liver was freeze-dried (2.5-L benchtop freeze-dry system; Labconco) for 48 h and fat was extracted with petroleum ether solvent (boiling point: 40–60°C) for 40 min using an ANKOM™2200 extractor (ANKOM Technology).

Blood analysis
Fasting plasma glucose, TG, total cholesterol, and total phosphorus were determined with an enzymatic colorimetric method on the Vitros Chemistry System (Johnson & Johnson Ortho-Clinical Diagnostics). The plasma insulin concentration was determined by an enzyme immunoassay using an insulin rat ELISA kit (EZRMI-13K; EMD Millipore Corporation).

Liver biopsy
Small liver sections were stained with hematoxylin and eosin (H&E) and Oil Red O (ORO) for evaluation of necroinflammatory grading and fatty droplets. Histologic changes were assessed by modifying the scoring system for grading and staging for nonalcoholic fatty liver disease as described by Kleiner et al. (27).
Histopathology examination
Rat liver tissue was processed into 3- to 4-μm-thick formalin-fixed paraffin-embedded tissue sections and stained with H&E. Histopathologic examination consisted of assessing a steatosis grade and distribution score of 0 (<5%), 1 (5–33%), 2 (33–66%), or 3 (>66%). Location was defined as a steatosis distribution score of 0 (zone 3), 1 (zone 1), 2 (azonal), or 3 (panacinar). Microvesicular steatosis was recorded as a score of either 0 (not present) or 1 (present). Lobular inflammation was semiquantified according to a score of 0 (<2 foci per ×200 field), 1 (2–4 foci per ×200 field), or 3 (>4 foci per ×200 field).

ORO examination
ORO staining was performed according to a previously described protocol (28). Briefly, fresh-frozen rat liver tissue was embedded into cryomolds and sectioned into 5-μm sections on a cryostat (Leica). Sections were then stained with ORO and were semiquantified with NIH ImageJ software (http://rsbweb.nih.gov/ij). Tissue sections were imaged at 5 high-power (×400) fields and converted to 8-bit grayscale images. This was compared against a predefined image threshold according to a rat liver section negative for steatosis and microvesicular steatosis on H&E and ORO staining, and image analysis for ORO surface area staining determined.

Statistical analysis
The required number of rats (n = 9) was based on previous weight gain data (6.0 ± 0.95 g/d), assuming a 25% difference in the mean with 90% statistical power and 5% significance. Data are expressed as means ± SDs for all values. Data analysis was performed using SPSS 23 software (IBM SPSS). Results were analyzed by one-factor ANOVA, and specific comparisons were made between each of the 5 groups using Fisher’s pairwise comparisons. P < 0.05 was considered significant.

Results
Weight gain, food intake, and energy efficiency
Although initial body weights were similar among the groups, the final body weights were significantly different. The final body weight of the control group (NP-0.3P) was significantly greater than that of the LP-0.015P and LP-0.056P groups. Among the low-protein groups, body weight increased with an increased dietary phosphorus content and the body weight of the LP-0.1P and LP-0.3P groups was not significantly different from that of the NP-0.3P group (Table 1).

The addition of phosphorus to low-protein diets increased weight gain (expressed as g/d); weight gain in the LP-0.1P and LP-0.3P groups was close to that of the NP-0.3P group (Figure 1A). Moreover, food intake was improved by increasing the phosphorus content of the low-protein diets. No difference in food intake was observed among the LP-0.1P, LP-0.3P, and NP-0.3P groups (Figure 1B, Suppplmental Figure 1). The difference in food intake of the low phosphorus (LP-0.015P and LP-0.056P) groups started with the introduction of the various diets and persisted throughout the experimental period (Supplemental Figure 1). The pattern of variation in energy efficiency [expressed as weight gain (g)/100 kcal] paralleled that of weight gain, in which energy efficiency of the low-protein groups increased with the addition of phosphorus to the diet, in which that of the LP-0.015P was lower than that of LP-0.056P that was lower than that of LP-0.1P. However, no differences were observed between energy efficiency of the LP-0.1P and LP-0.3P groups that was similar to that of the NP-0.3P group (Figure 1C).

TABLE 1  Body weight and body composition of rats fed a control diet or 1 of 4 low-protein diets with different phosphorus concentrations for 9 wk1

| Variable | LP-0.015P (n = 9) | LP-0.056P (n = 10) | LP-0.1P (n = 10) | LP-0.3P (n = 10) | NP-0.3P (n = 10) | ANOVA P value |
|----------|-----------------|-----------------|-----------------|-----------------|-----------------|--------------|
| Weight, g | 266.1 ± 35.7 | 265.4 ± 32.8 | 266.5 ± 27.6 | 267.8 ± 24.2 | 266.9 ± 31.9 | 1.000 |
| Water, % | 65.4 ± 2.1 | 62.4 ± 3.7 | 57.2 ± 6.3 | 53.9 ± 9.7 | 55.1 ± 5.6 | <0.001 |
| Fat, % | 24.9 ± 9.4 | 33.8 ± 15.8 | 56.6 ± 32.1 | 64.9 ± 37.7 | 56.4 ± 21.2 | <0.001 |
| Defatted | 20.6 ± 5.0 | 23.1 ± 6.1 | 27.3 ± 8.6 | 26.7 ± 8.5 | 24.2 ± 5.3 | 0.006 |
| Liver, % | 14.0 ± 3.7 | 14.5 ± 3.9 | 15.5 ± 5.6 | 19.5 ± 5.8 | 20.8 ± 5.4 | 0.010 |

1Values are means ± SDs. One-factor ANOVA was used to detect significant differences between groups. Significance was set at P < 0.05. Categories in the same row that do not share the same superscript letters are significantly different. LP-0.015P, 10% protein and 0.015% P; LP-0.056P, 10% protein and 0.056% P; LP-0.1P, 10% protein and 0.1% P; LP-0.3P, 10% protein and 0.3% P; NP-0.3P, 20% protein and 0.3% P (control group).
Body and liver composition

Results of the body proximate analysis suggested that the addition of phosphorus to low-protein diets was associated with an increase in body moisture. However, the proportion of body moisture decreased with the addition of phosphorus to low-protein diets. The proportion of moisture in the NP-0.3P group was similar to that of the LP-0.3P group and both were significantly less than the proportion of moisture in LP-0.015P– and LP-0.056P–fed rats. The addition of phosphorus to low-protein diets was associated with a significant increase in total body fat content ($P < 0.01$), but no

![Graphs showing weight gain, food intake, and energy efficiency against dietary phosphorus concentration.](https://academic.oup.com/cdn/article-abstract/1/8/e000943/4735235)
Table 2: Plasma metabolites of rats fed a control diet or 1 of 4 low-protein diets with different phosphorus concentrations for 9 wk

| Variable          | LP-0.015P (n = 9) | LP-0.056P (n = 10) | LP-0.1P (n = 10) | LP-0.3P (n = 10) | NP-0.3P (n = 10) | ANOVA P value |
|-------------------|------------------|------------------|------------------|------------------|------------------|---------------|
| Phosphorus, mg/dL | 5.4 ± 0.9a       | 6.5 ± 1.2b       | 7.4 ± 0.9b       | 6.4 ± 0.7a,b     | 6.4 ± 0.9b,b     | 0.002         |
| Glucose, mg/dL    | 148.3 ± 91.7a    | 205.0 ± 57.5b    | 245.4 ± 83.3b    | 263.9 ± 68.6b    | 277.0 ± 76.5b    | 0.004         |
| Insulin, ng/mL    | 0.4 ± 0.2        | 0.5 ± 0.4        | 0.8 ± 0.8        | 0.8 ± 0.9        | 0.7 ± 0.9        | 0.487         |
| TG, mg/dL         | 35.7 ± 6.7       | 41.2 ± 17.2      | 45.0 ± 22.7      | 40.6 ± 15.8      | 41.2 ± 13.1      | 0.807         |
| Cholesterol, mg/dL| 87.0 ± 22.1      | 85.3 ± 13.3      | 84.8 ± 14.7      | 85.5 ± 15.9      | 105.4 ± 25.3     | 0.081         |
| Albumin, mg/dL    | 3.2 ± 0.2        | 3.2 ± 0.3        | 3.1 ± 0.2        | 3.1 ± 0.2        | 3.2 ± 0.3        | 0.921         |
| C-reactive protein, mg/dL | 3.2 ± 0.4 | 2.8 ± 0.4        | 2.7 ± 0.7        | 2.7 ± 0.8        | 3.1 ± 0.3        | 0.162         |

1 Values are means ± SDs. Significance was set at P < 0.05. Categories in the same row that do not share the same superscript letters are significantly different. LP-0.015P, 10% protein and 0.015% P; LP-0.056P, 10% protein and 0.056% P; LP-0.1P, 10% protein and 0.1% P; LP-0.3P, 10% protein and 0.3% P; NP-0.3P, 20% protein and 0.3% P (control group).

significant changes in percentage of body fat were observed (P = 0.23). The defatted carcass weight was used to provide information on body protein status, because mineral content is known to be relatively small. The quantity and percentage of defatted rat carcasses in the LP-0.3P and NP-0.3P treatment groups was similar regardless of dietary protein content and was significantly greater than that of the LP-0.015P and LP-0.056P groups (Table 1). In the low-protein groups, liver weight increased with increasing dietary phosphorus but liver weights of LP-0.1P and LP-0.3P rats were similar to NP-0.3P rats. However, when liver weight was expressed per 100 g of body weight, no statistically significant differences were detected among treatment groups (P = 0.55). In addition, no significant differences in percentages of hepatic dry weight or fat content were observed among the groups (Table 1).

Plasma results

The total plasma phosphorus concentration was significantly different among groups (P < 0.01); the LP-0.015P treatment group had the lowest value, whereas the other treatment groups maintained similar levels. The plasma glucose concentration in the NP-0.3P group was similar to the LP-0.3P group. Among the low-protein groups, the plasma glucose concentration increased with the increased phosphorus content of the diet. Plasma TG, total cholesterol, albumin, and C-reactive protein concentrations were not significantly different among treatment groups and were not affected by either the protein or phosphorus content of the diets (Table 2).

In the low-protein treatment groups, plasma urea nitrogen decreased when the phosphorus content of the diet increased to ≤0.1% P. No differences in plasma urea nitrogen were observed among the NP-0.3P, LP-0.1P, and LP-0.3P groups (Figure 2).

Liver histology

Analyses of liver photomicrographs (Supplemental Figure 2) indicated that steatosis grade and location, microvesicular steatosis, and portal inflammation were similar among groups and thus were not affected by the protein or phosphorus content of the diet. All groups had approximately similar cases of microvesicular steatosis. Approximately 5–6 rats from each group presented with grade 0 or grade 1 steatosis, and only 1 rat from each group developed grade 3 steatosis. No signs of Mallory hyaline bodies, fibrosis, or glycogenated nuclei were detected in all rats. Only lobular inflammation was significantly greater in the low-protein groups than in the normal dietary protein group (Table 3).

Discussion

This study was designed to investigate the impact of dietary phosphorus supplementation on several growth parameters of rats maintained with a low-protein diet. In animals, food intake was regulated to serve 2 objectives. First, satisfying nutritional needs for growth and maintenance entails long-term regulation of intake, which is usually accompanied by an increase in food intake. In support of this, diet selection and protein restriction studies show that young animals are capable of adjusting their intake to support their nutritional protein requirements (29, 30) for growth and maintenance. Second, maintenance of homeostasis relates to acute or short-term regulation of food intake (31), which is usually associated with a...
decrease in food intake to avoid the toxicity of amino acid accumulation that is common with the ingestion of low-quality proteins. Thus, reduced food intake of the low-protein low-phosphorus diet groups (LP-0.015P and LP-0.056P) compared with the other low-protein groups (LP-0.1P and LP-0.3P), despite the similarity in dietary protein contents, is likely to have resulted from a dietary adaptation for the maintenance of homeostasis (32). Adequate protein metabolism in terms of synthesis and degradation depends on the availability of essential amino acids and energy needed to catalyze these reactions. Protein synthesis is a high energy-requiring process, in which 4 ATP equivalents are required for the formation of 1 peptide bond, or the equivalent of 0.67 kcal/1 g of protein synthesized (30). Several observations indicate that this process is highly dependent on phosphorus availability (35–37). In mice, dietary phosphorus restriction was reported to lower gastrocnemius body protein metabolism because the percentage of body protein (defatted percentage) of the 0.3% P (LP-0.3P and NP-0.3P) groups was higher than that of the LP-0.1P group. This further confirms the importance of phosphorus in protein metabolism and indicates that phosphorus is capable of altering body composition. In this study, no significant alterations in liver composition and histology were detected, despite the fact that dietary phosphorus restriction in mice is reported to increase hepatic lipid accumulation, especially under conditions of increased cholesterol content in the diet (41).

In addition, the high resemblance between the LP-0.1P and control (LP-0.3P and NP-0.3P) groups in terms of weight gain, food intake, and body weight was not reflected by similarities in body composition, because the percentage of body protein (defatted percentage) of the 0.3% P (LP-0.3P and NP-0.3P) groups was higher than that of the LP-0.1P group. This further confirms the importance of phosphorus in protein metabolism and indicates that phosphorus is capable of altering body composition. In this study, no significant alterations in liver composition and histology were detected, despite the fact that dietary phosphorus restriction in mice is reported to increase hepatic lipid accumulation, especially under conditions of increased cholesterol content in the diet (42).

Moreover, the failure of dietary phosphorus restriction to significantly affect plasma phosphorus content except under extreme conditions of restriction (LP-0.015P) is in line with findings in humans (43), in which plasma phosphorus is reported to not be an indicator of phosphorus intake. However, manipulation of dietary phosphorus (between 0.056% and 0.3% P) affected food intake, energy efficiency, weight gain, and body composition, although plasma phosphorus was not affected. This further confirms that

| Variable                          | LP-0.015P (n = 9) | LP-0.056P (n = 10) | LP-0.1P (n = 10) | LP-0.3P (n = 10) | NP-0.3P (n = 10) | ANOVA P value |
|-----------------------------------|------------------|-------------------|-----------------|-----------------|-----------------|--------------|
| Oil Red O image analysis, µm²     | 280 ± 727        | 1510 ± 2837       | 1234 ± 1636     | 2075 ± 3088     | 664 ± 1457      | 0.407        |
| Steatosis grade                   | 0.4 ± 1.0        | 0.5 ± 1.0         | 0.9 ± 1.1       | 0.6 ± 1.0       | 0.3 ± 0.7       | 0.701        |
| Location                          | 0.7 ± 1.0        | 0.7 ± 1.3         | 1.8 ± 1.6       | 0.9 ± 1.5       | 0.3 ± 0.9       | 0.118        |
| Microvesicular steatosis          | 0.6 ± 0.5        | 0.6 ± 0.5         | 0.9 ± 0.3       | 0.7 ± 0.5       | 0.5 ± 0.5       | 0.385        |
| Lobular inflammation              | 0.7 ± 0.9abc     | 0.7 ± 1.0abc      | 1.4 ± 1.1ab     | 0.5 ± 0.7ab     | 0.1 ± 0.3b      | 0.020        |
| Portal inflammation              | 0.2 ± 0.4        | 0.2 ± 0.4         | 0.1 ± 0.3       | 0.1 ± 0.3       | 0.1 ± 0.3       | 0.895        |

1Values are means ± SDs. One-factor ANOVA was used to detect significant differences between groups. Significance was set at P < 0.05. Categories in the same row that do not share the same subscripts are significantly different. LP-0.015P, 10% protein and 0.015% P; LP-0.056P, 10% protein and 0.056% P; LP-0.1P, 10% protein and 0.1% P; LP-0.3P, 10% protein and 0.3% P; NP-0.3P, 20% protein and 0.3% P (control group).
the plasma phosphorus concentration is neither a good marker of phosphorus intake nor an indicator of changes in food intake or weight measures.

Furthermore, the elevated plasma glucose observed in the control group may have been related to the effect of egg white protein on insulin secretion (44–46). Human studies have shown that the incremental insulin area was much lower in both healthy subjects (45) and individuals with diabetes (44) after ingestion of 50 g of egg white compared with cottage cheese. In addition, ingestion of a breakfast with whole egg, egg white, or egg yolk was associated with an increase in blood glucose concentration after egg white ingestion than after meals with whole egg or egg yolk (46). However, the resemblance in plasma glucose between LP-0.3P and NP-0.3P implies that plasma glucose is not solely dependent on the quantity of egg white in the diet. In low-protein (egg white) groups, the increase in plasma glucose with the addition of phosphorus possibly indicates that egg white ingestion is able to blunt the effect of phosphorus on insulin sensitivity (47) or the presence of varied glycogen storage between the groups. The latter option may prevail, because plasma glucose is affected by hepatic glycogen content that is known to be stimulated by phosphorus intake (48).

The ability of phosphorus intake to manipulate food intake and body composition requires further investigations to determine the level of phosphorus intake that is capable of improving body composition.

Low body protein content (defatted percentage) was present under conditions of 0.1% P (LP-0.1P; ~0.25 mg P/kcal) and was lower despite normal protein intake, whereas higher levels of phosphorus intake improved body protein content. The level of phosphorus intake (0.25 mg/kcal of highly bioavailable phosphorus) resembles that consumed by most people, which is ~0.5 mg P/kcal (49) assuming a 50% bioavailability (owing to the significant contribution of plant sources). This level is not thought to enhance protein status. In support of this, a recent study in humans reported an improvement in body weight and waist circumference of overweight and obese subjects after 12 wk of phosphorus supplementation (50). Although body composition was not measured, an increase in the percentage of lean body mass would be expected.

Under moderate protein restriction (10%), the addition of phosphorus (0.3%) was able to significantly improve food intake, weight gain, and energy efficiency similar to a normal protein diet (20%) containing the same concentration of phosphorus (0.3%). Body protein content (defatted percentage) was high with a 0.3% P diet, irrespective of protein content (10% or 20%). The pattern of changes in the different measures seems to indicate that a dietary phosphorus content of 0.1% may represent a critical threshold. However, it is not clear whether phosphorus would be able to exert these effects under added protein restriction. Our results suggest that not only the protein quantity but also the phosphorus content of the diet is of extreme importance for improving food intake, weight gain, and energy efficiency.

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