Factors affecting circulating levels of peptide YY in humans: a comprehensive review

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Abstract

As obesity continues to be a global epidemic, research into the mechanisms of hunger and satiety and how those signals act to regulate energy homeostasis persists. Peptide YY (PYY) is an acute satiety signal released upon nutrient ingestion and has been shown to decrease food intake when administered exogenously. More recently, investigators have studied how different factors influence PYY release and circulating levels in humans. Some of these factors include exercise, macronutrient composition of the diet, body-weight status, adiposity levels, sex, race and ageing. The present article provides a succinct and comprehensive review of the recent literature published on the different factors that influence PYY release and circulating levels in humans. Where human data are insufficient, evidence in animal or cell models is summarised. Additionally, the present review explores the recent findings on PYY responses to different dietary fatty acids and how this new line of research will make an impact on future studies on PYY. Human demographics, such as sex and age, do not appear to influence PYY levels. Conversely, adiposity or BMI, race and acute exercise all influence circulating PYY levels. Both dietary fat and protein strongly stimulate PYY release. Furthermore, MUFA appear to result in a smaller PYY response compared with SFA and PUFA. PYY levels appear to be affected by acute exercise, macronutrient composition, adiposity, race and the composition of fatty acids from dietary fat.

Key words: Satiety; Exercise; Dietary fatty acids; Obesity

Introduction

The prevalence of obesity in the USA continues to be a public health concern, with approximately two-thirds of the adult population being classified as overweight and 35% being classified as obese[1,2]. While these rates are highest in the USA, the UK and all of Europe also have a high prevalence of adults being classified as overweight (range 38–60%) or obese (approximately 20–25% of adults)[3,4]. Obesity is associated with hypertension, dyslipidaemia, CVD, stroke, diabetes mellitus, musculoskeletal disorders, certain cancers and an increased risk of disability. Thus, obese individuals have an increased risk for morbidity and mortality[5,6]. The growing epidemic of obesity is attributed primarily to lifestyle factors, specifically by decreases in physical activity, increases in total energy and fat intake, and the interaction of these environmental factors with genetic susceptibility[7].

To maintain body weight, average daily energy intake must be matched with average daily energy expenditure so that an individual is in energy balance. Only when energy balance is achieved can the prevention of future weight gain occur. Although meal-to-meal and day-to-day variations are quite large, over time, energy intake is matched to energy expenditure and body-weight changes are minimal or modest over periods of months and years[8]. Consequently, it is thought that energy intake must also be well regulated. Not all research supports this idea, though, and it has recently been an issue of debate among scientists, with some supporting the notion that energy balance is not tightly regulated[9,10]. One measure of support for this long-term regulation over energy balance is the history of weight gain in adults. Average annual weight gain since the early 1960s (which was the beginning of the obesity epidemic) is a modest 0·3 kg[11,12]. This equates to an energy surplus of about 29 kJ/d (7 kcal/d). This is, of course, without correcting for a greater energy expenditure as body weight increases which would result in a larger energy surplus if assessed over a period of decades where substantial weight gain is occurring[9]. Regardless of the amount of energy surplus, it is also important to remember that several factors aside from satiety hormones affect what

Abbreviations: ARC, arcuate nucleus; CCK, cholecystokinin; NPY, neuropeptide Y; POMC, pro-opiomelanocortin; PYY, peptide YY.

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we eat including emotions, social factors, time of day, convenience and cost. During the past 20 years, research into the mechanisms of hunger and satiety has identified numerous circulating hormones that are thought to play a role in the overall control of hunger and satiety and thus act to regulate total energy intake, and to a lesser extent energy expenditure, which ultimately affects body weight. One such satiety hormone, peptide YY (PYY), has been assessed recently in a number of studies that are seeking to determine what nutrients, human demographics, or type of physical activity make an impact on circulating PYY levels to the greatest extent. There are two purposes of the present review: (1) to summarise the recent literature on the role of PYY in regulating energy balance and the factors that affect its release and circulating levels within the human body; and (2) to examine the emerging evidence on the effect of dietary fatty acid composition on PYY release and circulating levels in humans.

Discussion

Structure of peptide YY

PYY was first identified in the small intestine by Tatemoto & Mutt in 1980. It is a thirty-six-amino acid peptide that has both an amino acid terminal tyrosine and a carboxyl terminal tyrosine amide and belongs to the pancreatic polypeptide family. Its tertiary structure consists of a polyproline helix and α-helix connected by a β-turn resulting in a characteristic U-shaped peptide. PYY is secreted mostly from the endocrine L-cells of the ileum, colon and rectum (with highest concentrations in the colon and rectum), although PYY mRNA has been identified in the stomach, duodenum, jejunum, pancreas and brainstem. Its secretion is initiated either by direct luminal contact of nutrients with the endocrine cells or indirectly through neurohumoral signals. These neurohumoral signals are hormones that are released into the blood by neuroendocrine cells that are stimulated through nerve impulses. PYY exists in two forms: PYY1–36 and PYY3–36. PYY5–36 is produced by the enzyme dipeptidyl peptidase-IV which hydrolyses PYY at the Pro2–Ile3 bond of the NH2 terminus. PYY1–36 binds and activates all five Y receptors (Y1–Y5). If the first two amino acids at the N-terminal are removed, the receptor selectivity changes because PYY5–36 only has a high affinity for the Y2 receptor. While both forms are biologically active at all times, PYY3–36 is the main storage and circulating form of PYY and is thought to more actively control food intake.

Functions of peptide YY

PYY has multiple functions including slowing gastric emptying and gastrointestinal motility, inhibiting the secretion of gastric acid, gallbladder contractions, and pancreatic exocrine enzymes, and the most well-documented function of regulating food intake. Intravenous infusions of PYY reduce appetite and energy intake in healthy human subjects, suggesting that PYY plays a role in regulating satiety. Circulating levels of PYY are low in the fasted state and both the 3–36 and 1–36 forms increase rapidly upon nutrient ingestion. While it is well known that PYY is released in response to nutrient intake, it is also released by gastric acid, cholecystokinin (CCK), and infusion of bile acids into the ileum or colon as shown in animal studies. Finally, PYY secretion is affected by things other than food consumption such as intestinal peristalsis and intraluminal nutrients. Much like ghrelin and CCK, PYY is not released by gastric distension.

Plasma PYY levels increase within 15 min after nutrient consumption, peak at about 60–90 min, and remain elevated for up to 6 h. Postprandial levels have been shown to be proportional to meal size. The initial increase in PYY concentration occurs before food is able to reach the L-cells of the small intestine, indicating that there is a neural or endocrine mechanism that involves cephalic phase-mediated PYY release. PYY release becomes greater when the nutrients arrive at the endocrine L-cells and the sustained release of PYY for up to 6 h postprandially is thought to be due to the direct effects of the intraluminal gut contents on the L-cells. PYY responds to all three macronutrients, with the strongest stimulus being with either fats or proteins (described in more detail below). Appetite-suppressant effects of duodenal lipids are dependent on fat digestion, so fat digestion may also be a prerequisite for fat-induced stimulation of PYY. Indeed, blocking the hydrolysis of long-chain fatty acids abolishes the typical PYY postprandial response, probably due to a lack of CCK release.

Other foods or substances have also been tested for potential PYY responses. PYY has been shown to increase following ingestion of decaffeinated coffee, but does not change in response to alcohol ingestion in adult men and women. Additionally, it has recently been suggested that artificial sweeteners could stimulate the release of PYY because of two sweet taste receptors (T1r2 and T1r3) that are co-localised with PYY in enteroendocrine L-cells. However, ingestion of the artificial sweetener sucralose does not lead to changes in circulating PYY levels.

Mechanism of action for peptide YY

Similar to other peripheral hunger or satiety signals, PYY acts on the arcuate nucleus (ARC) of the hypothalamus by targeting the neuropeptide Y (NPY) neurons. PYY is able to cross the blood–brain barrier freely by a non-saturable mechanism. This non-saturable mechanism implies that there is no point at which saturation of that mechanism occurs, so there is no limit to the amount of
PYY that can cross that blood–brain barrier. Within the ARC, there are two subsets of neurons that integrate signals and influence energy homeostasis. These neurons are the NPY/agouti-related peptide (AgRP) neurons and the pro-opiomelanocortin (POMC)/cocaine- and amphetamine-regulated transcript (CART) neurons(40,41). The NPY/AgRP neurons are classified as orexigenic (appetite stimulating) while the POMC/CART neurons are classified as anorexigenic (appetite suppressing)(42). In the ARC, PYY is a potent agonist of both the Y1 and Y2 receptors and can also bind to the Y5 receptor whereas PYY3–36 has a very high affinity for the Y2 receptor(37). The Y2 receptor of NPY is a 381-amino acid, seven transmembrane-spanning, G-protein-coupled receptor(43). In humans, the Y2 receptor mRNA which includes the nucleus tractus solitarius(19). An inhibition of food intake(25). Further, Y2 receptor knock-out mice are unable to inhibit food intake in the presence of PYY3–36. Additionally, in rodents, injection of PYY3–36 directly into the ARC inhibits release of NPY, stimulates release of α-MSH, and inhibits food intake(19,24). In another rodent study, PYY3–36 administered at a dose that is representative of peak postprandial plasma levels resulted in decreased food intake during the dark phase (the dark phase is when rodents are awake, so they are moving around and eating during this phase). The rodents also received twice-daily doses of PYY3–36 for 8 d, which resulted in a significant reduction in food intake and reduced body weight when compared with the saline control(19).

The satiating effects of PYY have been clearly shown in human subjects as well. Batterham et al.(24) gave a 90-min infusion of PYY3–36 to twelve normal-weight subjects (six men and six women). The dose was designed to mimic normal postprandial responses. They reported a decreased energy intake of 36% at a free buffet meal following PYY administration. Further, food diaries showed that food intake remained decreased for up to 12 h following a PYY infusion, but did not alter food intake from 12 to 24 h after administration. The total reduction in energy intake over a 24 h period was 33%(24). Exogenous administration of PYY was also used to study its effects in lean and obese adults. Energy intake at an ad libitum buffet 2 h after PYY3–36 infusions was decreased by 31% in normal-weight adults and 30% in obese adults(25). The aforementioned findings for a reduction in food intake following exogenous administration of PYY in both normal-weight and obese adults have been confirmed in several other studies(47–50).

Together, these studies provide further evidence of the acute or short-term regulation of PYY3–36 on energy intake in both human and rodent models.

**Role of peptide YY in appetite control**

While the aforementioned paragraphs provide clear evidence on the response of PYY to nutrient ingestion, as well as the impact that exogenously administered PYY has on decreasing food intake and body weight, the role of endogenously produced PYY in controlling overall appetite is less frequently studied and less conclusive. Some of the controversy over the role of endogenous PYY on appetite control may stem from some reports that PYY levels are not always correlated with subjective
ratings of hunger and fullness. Importantly, though, there is also evidence that visual analogue scales may not be a good indicator of subjective feelings of hunger and fullness. This limits the ability to determine whether or not PYY directly affects subjective feelings of hunger and fullness. To further complicate the issue, other studies have shown that plasma PYY levels are positively correlated with ratings of fullness and inversely correlated with ratings of hunger.

The other area of research that generates questions about the role of endogenous PYY levels on appetite control is the relationship between circulating PYY levels and daily food intake or intake at an ad libitum buffet. In rodents, circulating PYY levels following test meals of varying protein content were inversely correlated with food intake. This suggests at least a partial role of PYY in appetite regulation. Conversely, in human subjects, neither fasting nor postprandial PYY levels were associated with appetite regulation. Conversely, in human subjects, neither fasting nor postprandial PYY levels were associated with appetite regulation. This limits the ability to determine whether or not PYY directly affects subjective feelings of hunger and fullness. To further complicate the issue, other studies have shown that plasma PYY levels are positively correlated with ratings of fullness and inversely correlated with ratings of hunger.

Peptide YY levels in non-obese v. obese

Fasting PYY levels correlate negatively with BMI, as levels in obese individuals are lower in the fasting state compared with non-obese individuals. This has been shown in both adults and children. Further, secretion of PYY following a meal has also been shown to be lower in obese subjects as compared with lean subjects. This holds true even if obese subjects consume more food from a buffet meal than lean counterparts. It has also been shown that weight loss leads to an increase in PYY, suggesting that weight loss can restore PYY levels. Combined, these studies imply that there may be a PYY deficiency in the pathogenesis of obesity or that an altered PYY phenotype exists following the onset of obesity. However, obesity does not appear to be associated with PYY resistance as is the case for leptin. PYY infusions in obese subjects result in a similar percentage reduction in food intake as compared with lean subjects. The mechanism(s) by which obese individuals have an altered PYY profile remain unknown. It could be the result of abnormalities in its synthesis, release, and/or clearance; however, more work is needed to determine the causes for altered PYY fasting and postprandial profiles in obese individuals.

To complicate this association between weight status and PYY levels, some studies have shown no relationship between BMI or body weight and PYY levels. It has been reported that weight loss in obese adolescents did not lead to changes in fasting PYY levels. Kim et al. also reported that fasting PYY was not correlated with body weight, and PYY AUC was not different between lean and obese older adults following an oral glucose tolerance test. While it is unknown what causes the discrepancies between studies on the relationship between fasting PYY and BMI, the type of test meal incorporated in studies may have an impact on postprandial responses. In studies where differences were seen between obese and lean participants, mixed meals were used. Conversely, the study by Kim et al. where there were no differences in PYY based on body weight, used an oral glucose tolerance test. It has also been shown that a greater postprandial PYY for normal-weight v. obese subjects exists for high-fat test meals, but not for high-protein or high-carbohydrate test meals. The presence of dietary fat may be necessary to show differences in postprandial PYY levels based on weight status. It has been shown that fat components directly stimulate L-cell secretion through activation of G-protein coupled receptor (GPR)-120 receptors. It is possible that in obese individuals, GPR-120 receptors are down-regulated to fat stimulation, which would decrease PYY secretion.

Interestingly, one study has even reported that PYY levels were positively associated with BMI and waist circumference. This statistically significant correlation was weak (r=0.158), and when participants were put into tertiles based on PYY levels, the BMI levels between the tertiles were quite small (range 27.8–29.5 kg/m²). Clearly more research is needed to fully understand the relationship between body-weight status and both fasting and postprandial PYY levels. Additionally, although there
appears to be some effect of body-weight status on PYY levels, it remains to be shown what the mechanism is behind that effect. There may be a genetic component, since plasma PYY was shown to be highly heritable in a phenotyped twin sample. A genetic pleiotropy was also found between PYY levels and BMI, suggesting that genetic variation at the PYY locus may influence weight status and other heritable metabolic syndrome traits.

Peptide YY as a therapeutic target for obesity

Because of its well-known role in appetite regulation, PYY analogues are a therapeutic target for weight loss and weight management. Many initially thought that exogenous PYY administration could lead to a potential cure for the obesity epidemic. Indeed, exogenous administration of PYY$_{3-36}$ has been shown to reduce body weight and adiposity in normal-weight and obese rodents fed a high-fat diet. Additionally, PYY transgenic mice that are crossed with ob/ob (genetically obese) mice exhibit lower body weights and adiposity. There are currently two main issues decreasing the potential for PYY analogues to be effective as a therapeutic target for weight loss. The first issue is the severe gastrointestinal side effects that have been reported with moderate doses up to supraphysiological doses of PYY$_{3-36}$, which greatly limits its impact as a therapeutic target for obesity. The other issue deals with the method of administration. Oral ingestion of the hormone is not absorbed well and some argue that subcutaneous injections are not as effective because they are not directly stimulating the L-cells of the gastrointestinal tract. More recently reported successful absorption of oral administration of PYY$_{3-36}$ with sodium N-caprylate in adult human subjects. Unfortunately, acute and 24 h energy intake was unchanged by the PYY administration, which again limits its potential as a therapeutic target to treat obesity. Therefore, slow progress is being made on the therapeutic front, but many roadblocks still remain. It is also important to recognise that overall energy homeostasis is influenced by a number of parameters and one single hormone is unlikely to be sufficient to permanently alter energy balance.

Effects of exercise on peptide YY

In the past 3 to 4 years, a number of studies have emerged assessing PYY responses to exercise. We, and others, have shown that acute aerobic exercise (ranging from low to high intensity) results in higher circulating plasma PYY levels. While one recent study found no differences in post-exercise PYY for either continuous aerobic exercise or intermittent exercise compared with sedentary conditions, most studies show acute increases in PYY levels. Further, this increase appears to be intensity dependent, as greater exercise intensity leads to greater circulating PYY levels. One study showed that steady-state and high-intensity intermittent exercise both raised post-exercise PYY; however, PYY in the hours following exercise were highest following the high-intensity intermittent exercise. Resistance exercise also acutely increases PYY levels, albeit in a lesser magnitude than aerobic exercise of similar energy output. Further, subjective scores of hunger during both aerobic and resistance exercise are reduced. The increase in PYY following exercise is only an acute response. There is no chronic adaptation to long-term exercise training. This was shown by the lack of change in fasting PYY levels after both aerobic training and resistance training regimens.

Increased PYY levels as the result of aerobic exercise have also been shown to reduce subsequent energy intake in both lean and obese individuals, indicating an even greater protective effect against weight gain. Still other studies have shown no differences in energy intake (or even slight increases in energy intake) when comparing post-exercise vs. sedentary conditions. Importantly, however, if one takes into account the energy expended during exercise, there is still a greater negative energy balance in those studies exhibiting greater post-exercise PYY levels.

A few studies have also examined the effects of chronic aerobic exercise. One challenge to these studies is the fact that weight loss usually occurs, making it difficult to determine whether changes in PYY are due to the exercise, weight loss, or a combination of the two. In obese adolescents, exercise training increased fasting PYY levels; however, some loss in fat mass also occurred which could at least partially explain the changes in PYY concentrations. Conversely, a study in obese adults who performed aerobic training for 15 d revealed no differences in fasting or postprandial PYY levels from before to after the training intervention. Yet another study showed no significant differences in PYY responses to an acute bout of exercise following a 12-week aerobic exercise programme. There was a trend ($P$=0.06) for higher PYY values post-training; however, significant loss of body weight also occurred which could give at least partially explain the changes in PYY concentrations. Probably the best study to elucidate the effects of chronic exercise independent of weight loss was done by Scheid et al. where they showed that fasting PYY did not change following a 3-month exercise intervention in the absence of weight loss. Therefore, it appears that the effect of exercise on PYY is an acute response and that independent of changes in body weight, PYY levels do not change with chronic exercise training.

Effects of ethnicity, sex and age on peptide YY levels

Since weight status appears to influence PYY levels, investigators have explored whether circulating levels of PYY,
either in the fasting or postprandial state, differ based on sex, ethnicity or age. To date, the research on race comparisons has been done in Caucasian and African-American individuals. One study in prepubertal children showed that fasting PYY, and PYY following an oral glucose tolerance test, were lower in African-American compared with Caucasian children\textsuperscript{(91)}. Similarly, studies in adults also point toward reduced PYY levels in African-American individuals. One study examining both weight status (obese \textit{v.} normal weight) and race reported no differences in fasting PYY levels between normal-weight or obese African-American \textit{v.} Caucasian women. However, postprandial PYY levels were lower in obese African-American women compared with normal-weight African-American women, and both normal-weight and obese Caucasian women\textsuperscript{(92)}. Finally, another study comparing African-American \textit{v.} Caucasian women (matched for age and BMI) showed that both fasting and postprandial PYY levels were significantly lower in the African-American compared with Caucasian groups. Further, the change in PYY from baseline to 8 h postprandial was also lower in the African-American subjects\textsuperscript{(93)}. Together, these studies suggest that the PYY meal response is blunted in African-American compared with Caucasian women and that fasting PYY levels may also be lower. Since African-American women have a higher rate of obesity compared with Caucasian women in the USA\textsuperscript{(94,95)}, understanding the reason behind these differences in PYY could make an impact on future therapeutic targets in African-American populations.

Several studies have explored the possibility of differences in fasting or postprandial PYY in humans based on sex, with conflicting results. One limitation in making comparisons between men and women is the fact that body weight and adiposity are generally not the same between the sexes and this may make an impact on differences in PYY, independent of sex. Indeed, one study showed that women had lower fasting and postprandial PYY levels\textsuperscript{(96)}. However, the male subjects also weighed more and the PYY levels were not adjusted for body weight. Therefore, the elevated PYY levels may be due to the larger body weights in men \textit{v.} women. Conversely, Kim \textit{et al.}\textsuperscript{(66)} found that while fasting PYY levels did not differ between men and women, postprandial PYY (AUC) after an oral glucose tolerance test was significantly higher in women compared with men. More recent studies have reported no differences in PYY based on sex. This lack of difference was reported for both resting and acute exercise conditions\textsuperscript{(97)}, fasting conditions in normal-weight adults, even without adjustment for differences in body weight or adiposity\textsuperscript{(98)}, and with fasting conditions in both lean and obese children\textsuperscript{(67)}. Therefore, it appears that fasting PYY levels probably do not differ between men and women; however, there may be differences in acute meal responses. Whether postprandial PYY is higher or lower in men \textit{v.} women remains to be determined and could be affected by the type of meal consumed as well as the weight status or amount of adiposity in the individuals.

Finally, ageing has been associated with a decrease in appetite\textsuperscript{(39)} and a slowing of gastric emptying\textsuperscript{(100)}. Since PYY may influence both of these, it was plausible to expect that there may be a role of ageing on circulating PYY levels and responses to nutrients. While CCK has been reported to be higher in older \textit{v.} younger adults receiving an intraduodenal infusion of lipids and glucose, PYY does not appear to be affected by age\textsuperscript{(63,101)}.

**Impact of macronutrients on peptide YY**

Postprandial PYY concentrations have been shown to be influenced by the macronutrient content of the meal. Unfortunately, at this time, the macronutrient resulting in the greatest PYY response is still being debated by some because it may be influenced by meal composition, obesity status, sex, and possibly age. Nearly all studies are in agreement that carbohydrates induce the smallest PYY response. The debate is whether dietary fats or dietary proteins elicit the greatest PYY response. Early research showed that dietary fat elicited the greatest PYY response\textsuperscript{(18)} while more recent studies have shown that protein results in the greatest PYY response\textsuperscript{(55)}. It does appear that more studies support protein as a greater stimulator of PYY levels postprandially compared with fat. This has been shown in normal-weight and obese adults as well as obese adolescent females\textsuperscript{(55,66)}. Importantly, subsequent energy intake and ratings of hunger are also lowest following a protein-rich meal\textsuperscript{(55,66)}. Another study in adolescents showed that the greatest and most sustained PYY response was found after a protein-rich meal followed by a fat-rich meal, and finally by a carbohydrate-rich meal in obese adolescents; however, the normal-weight adolescents responded most strongly to the fat-rich meal\textsuperscript{(102)}. Therefore, in that study, weight status influenced the magnitude of PYY response to each macronutrient.

As mentioned above, it appears that both fat and protein elicit a greater postprandial PYY response compared with carbohydrates. One study in obese women during energy restriction reported greater PYY levels from high-protein \textit{v.} high-carbohydrate meals. Unfortunately, this did not translate to differences in subjective hunger or fullness responses\textsuperscript{(55)}. A single high-fat meal also resulted in greater postprandial PYY levels compared with a high-carbohydrate meal in obese women\textsuperscript{(103)}. Further, a 1-week diet that was either low in carbohydrates (high-fat) or high in carbohydrates (low-fat) led to a greater PYY response with the low-carbohydrate (high-fat) diet following test meals of identical composition\textsuperscript{(90)}. This suggests that diets or single meals rich in dietary fat or protein induce a greater PYY response compared with diets or meals rich in carbohydrates. As mentioned above, however, weight status, and possibly age, may affect this response. A study in obese and normal-weight
adolescent girls showed that normal-weight girls had a greater PYY response than obese girls to a high-fat meal while no differences between the groups were found following the high-carbohydrate meal. This again points towards the impact that weight status may play on the magnitude of PYY response to different macronutrients. Additionally, the use of different fatty acids or sources of protein could explain some of the discrepancies about which macronutrient stimulates the greatest PYY response. This is explored more in subsequent paragraphs. Finally, all of the aforementioned studies examined acute meal responses. It does not appear that a chronic high-fat diet leads to any differences in fasting PYY levels when compared with a low-fat (high-carbohydrate) diet.

It is also possible that the type of carbohydrate, fat or protein may influence the magnitude of PYY response. Very little research has been done on the type of protein and its effect on PYY release. Two studies in human subjects have reported no differences in the PYY response to a whey v. casein protein meal or whey v. both pea protein hydrolysate and a combined casein/whey milk protein (80% casein, 20% whey). Further, a 10-week feeding study in rats found no differences in plasma PYY levels between whey and soya protein diets. Conversely, PYY mRNA was down-regulated in rats following a lifetime of a high-whey diet v. high-casein or high-soya diets. Clearly, more studies are needed to determine the effects of different types of protein or amino acids on PYY levels in humans; however, initial work in human subjects and rats suggests no difference in circulating PYY levels based on different types of dietary protein.

With respect to carbohydrates, one study on glycaemic load and several studies on fibre have been published. The effect of a 1-week high-v. low-glycaemic-load diet was carried out in adult women. No effect was reported on either fasting or postprandial PYY levels. The effects of fibre on PYY have been examined in both acute and long-term diet studies. The addition of wheat fibre to a carbohydrate-rich test meal resulted in a blunted postprandial PYY response compared with control, whereas oat fibre did not differ from the control meal. Conversely, a pre-load of brown beans (higher in fibre) led to significantly higher fasting PYY levels and a greater PYY response at a test meal 10 h following the pre-load when compared with white wheat bread (lower in fibre). One long-term fibre supplementation study (4 g/d for 14 weeks) reported an increase in fasting PYY levels, while no differences in postprandial PYY levels were found. Similarly, 3 weeks of supplementing with 5–10 g of a fibre supplement per d led to higher fasting PYY levels in adult men and women. Therefore, chronic intake of a fibre supplement or a higher-fibre diet appears to increase fasting PYY levels while post-meal responses following either acute meals or chronic feeding studies remain controversial.

**Dietary fatty acids and peptide YY**

While PYY has been shown to respond to dietary fats, little to no research exists on whether or not the composition of fatty acids can differentially influence PYY release. Only in the last few years has some research in this area emerged. Investigators have examined the form of fat (NEFA v. TAG), the chain length of fatty acids and saturation of fatty acids. In normal-weight men, NEFA administered intra-gastrically resulted in greater PYY levels than TAG. With respect to chain length, intraduodenal infusions of either lauric acid (12 : 0) or capric acid (10 : 0) in healthy men showed that PYY levels were greater following 12 : 0 infusion compared with 10 : 0 and control, while 10 : 0 had no effect on PYY levels. Additionally, another study showed that intraduodenal infusions of oleic acid (18 : 1) led to a greater PYY response compared with lauric acid (12 : 0). Therefore, it appears that chain length is associated with PYY levels (the longer the chain length, the greater the PYY response).

With respect to fatty acid saturation, all of the studies have been done using long-chain fatty acids. Some previous research on subjective feelings of hunger and fullness indicate that fatty acids may influence overall hunger and satiety. Lawton et al. reported that a meal rich in PUFAs exerted the strongest control over appetite, based on subjective responses from visual analogue scale questionnaires as well as energy intake over the whole test day. Only four human studies to date have examined PYY responses to high-fat meals or high-fat diets rich in different fatty acids. These studies all examined fatty acid saturation, rather than chain length, on PYY responses. For the one high-fat diet study, we showed that a 50 high-fat diet rich in SFA resulted in higher postprandial PYY levels compared with a high-fat diet rich in MUFA in normal-weight men. This was only found after an evening meal, and 24 h measures of PYY did not differ between the diets. Additionally, these effects were found only in conjunction with daily aerobic exercise.

The other three remaining studies examined an acute or single high-fat meal response. Maljaars et al. reported no differences in PYY levels in normal-weight adults (men and women) following infusions of MUFA, PUFAs or SFA. It is important to note that these were infusions of fatty acids via a naso-ileal catheter and not oral consumption of foods containing those fatty acids. Conversely, Robertson et al. studied oral consumption of high-fat meals rich in either SFA, MUFA or PUFA. They showed a trend for both SFA- and PUFA-rich meals to stimulate a greater PYY response compared with a MUFA-rich meal in postmenopausal women. Finally, we recently showed that in normal-weight, premenopausal women, postprandial PYY levels were significantly higher for high-fatt meals rich in PUFA and SFA compared with MUFA. Based on the evidence published so far, it appears that the fatty acid composition of a meal or diet...
may influence the magnitude of postprandial PYY responses, and that PUFA and SFA induce a greater PYY response compared with MUFA. Since MUFA are considered more ‘heart healthy’ than SFA, this information could have significant implications for public health recommendations. If MUFA are promoted over SFA, it may be important to suggest including these fats with foods that may have a greater impact on satiety, such as lean sources of protein or fibre in order to prevent overconsumption.

**Mechanisms behind the role of fatty acid composition on peptide YY levels**

Since research on the significant effects of dietary fatty acid composition on postprandial PYY levels in human subjects has only recently been published, little to no information exists on the mechanisms behind these effects. Because PYY release is affected by CCK levels (29), it is possible that the effect of fatty acid composition on PYY is an indirect one through CCK. In the study mentioned above by Robertson et al. (122) it was shown that peak CCK release after a high-fat meal challenge was greatest with a meal rich in SFA compared with either MUFA or PUFA. Since CCK does influence PYY release, greater CCK following a SFA-rich meal may be at least one of the mechanisms behind the greater PYY levels that we reported for SFA- v. MUFA-rich high-fat meals (51).

Another hypothesis is related to the rate of digestion and absorption of different long-chain fatty acids. It has been shown in some very early studies that oils higher in MUFA and PUFA are absorbed more quickly than oils rich in SFA (123–126). While the present study was carried out in rodents, a human study showed support for slower absorption of SFA-rich high-fat meals compared with MUFA- or PUFA-rich meals based on postprandial chylomicron appearance in plasma (127). A slower absorption rate for SFA would mean that these fatty acids would be in contact with the endocrine L-cells that release PYY for longer periods of time. Therefore, the absorption rate of different fatty acids could probably affect the magnitude of PYY response. Since both of the aforementioned hypotheses have not been exclusively shown to be true, more work is needed to fully elucidate the mechanisms behind the differential PYY response to different dietary fatty acids. Additionally, more research is needed to see if the differences we reported in normal-weight, premenopausal women persist in other populations, such as males, obese individuals, and adolescents or children.

**Future direction**

Despite the fact that there are literally hundreds of studies that have been published on PYY, very little is known about the mechanisms by which nutrient ingestion stimulates the release of PYY from the enteroendocrine L-cells of the gastrointestinal tract. Some of the hypotheses mentioned above regarding the mechanisms behind different PYY responses to various dietary fatty acids may also be appropriate for other nutrients. It is known that CCK affects PYY release, although the exact mechanism behind this regulation is not clear (29). It has also been shown that dietary fat can directly stimulate PYY secretion from the L-cells through activation of GPR-120 receptors (67). Whether there are other proteins or hormones that can directly interact with the L-cells to release PYY remains to be determined. Additionally, we mentioned above that the length of time that nutrients are in contact with the L-cells influences the degree of PYY release; however, this has not been conclusively shown (51). Therefore, there is a need for more research to elucidate the mechanisms by which nutrients affect the release of PYY as well as the factors that influence the magnitude of PYY release independent of total energy consumption. Finally, our recent findings on the differential release of PYY in response to high-fat meals rich in different fatty acids opens up a new avenue for research. Previous studies have focused on how macronutrients (carbohydrates, fats and proteins) influence PYY response, with little thought as to the type of carbohydrate, fat or protein being tested. More research on the influence of different dietary sources or types of carbohydrates, fats or proteins on PYY levels is warranted.

**Conclusion**

Great progress in understanding the role of PYY as a satiety hormone and its influence on energy homeostasis has occurred in the last 30 years. Further, we now know more about the other actions of PYY in the body, how exercise and different foods or nutrients affect PYY levels, and how age, sex, ethnicity and weight status can make an impact on PYY levels. Since obesity continues to be an epidemic of great concern in the USA and worldwide, additional research on the possibilities of using PYY as a therapeutic target for obesity without the negative side effects is warranted. However, as mentioned above, there are challenges facing the therapeutic potential of PYY given that the common side effect of PYY administration is nausea and that there are a number of hunger and satiety signals that probably influence overall energy intake. Therefore, future studies should explore if chronic, low doses of PYY can be effective at reducing food intake over time without the negative side effects as well as the development of a way to administer PYY in order to bypass the negative gastrointestinal issues that currently occur. Future work on other pharmacological, nutrient and dietary options as a potential way to alter PYY levels, and possibly energy intake, should also be explored. Finally, the mechanisms behind the release of PYY following nutrient ingestion and how different types of fatty acids, proteins or carbohydrates can affect the magnitude of release need to be studied further.
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References

Peptide YY responses in humans


