Nuclear Magnetic Resonance Spectroscopy-Based Metabolomics of the Fatty Pancreas: Implicating Fat in Pancreatic Pathology

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Obesity • Inflammation • Metabolomics • Pancreatitis • Pancreatic cancer

Abstract
Background: Obesity is a worldwide epidemic and a significant risk factor for pancreatic diseases including pancreatitis and pancreatic cancer; the mechanisms underlying this association are unknown. Metabolomics is a powerful new analytical approach for describing the metabolome (complement of small molecules) of cells, tissue or biofluids at any given time. Our aim was to analyze pancreatic fat content in lean and congenitally obese mice using both metabolomic analysis and conventional chromatography.

Methods: The pancreatic fat content of 12 lean (C57BL/6J), 12 obese leptin-deficient (Lepob) and 12 obese hyperleptinemic (Lepdb) mice was evaluated by metabolomic analysis, thin-layer and gas chromatography.

Results: Pancreata of congenitally obese mice had significantly more total pancreatic fat, triglycerides and free fatty acids, but significantly less phospholipids and cholesterol than those of lean mice. Metabolomic analysis showed excellent correlation with thin-layer and gas chromatography in measuring total fat, triglycerides and phospholipids.

Conclusions: Differences in pancreatic fat content and character may have important implications when considering the local pancreatic proinflammatory milieu in obesity. Metabolomic analysis is a valid, powerful tool with which to further define the mechanisms by which fat impacts pancreatic disease.

Introduction

The burgeoning obesity epidemic in the United States has focused a spotlight on the role of adipose tissue in the multiple pathologic effects of obesity [1–4]. Adipose tissue secretes a number of proteins collectively referred to as adipokines that are important modulators of metabolism, inflammation and energy intake [5]. Obesity leads to dysfunctional production of adipokines. Specifically, circulating concentrations of the proinflammatory adipokine leptin are elevated, and those of the anti-inflammatory adipokine adiponectin are paradoxically de-
increased. This imbalance results in a generalized proinflammatory milieu, which is manifest at the local level by increased tissue macrophage infiltration as well as by increased production of the proinflammatory cytokines tumor necrosis factor-α (TNF-α), interleukin-6 (IL-6) and IL-1β [5, 6]. Fat infiltration of an organ is a known trigger for initiating this inflammatory cascade, with nonalcoholic steatohepatitis being a well-established example of this phenomenon [1, 7].

Clinical evidence also suggests that fat plays an important role in the development of pancreatic disease. Obesity is clearly a risk factor for increased severity of acute pancreatitis [8–11], and numerous well-controlled epidemiological studies have linked obesity to an increased incidence of pancreatic cancer [12–15]. In addition, some experimental work has begun to address the mechanisms by which fat contributes to pancreatic disease. Recent data from our laboratory showed that obese Leprob mice have an elevated pancreatic fat content and increased baseline levels of the proinflammatory cytokines TNF-α and IL-1β relative to lean wild-type mice [3]. In addition, obese leptin-deficient Lepob and obese hyperleptinemic Lepdb mice develop more severe pancreatitis than lean mice when subjected to cerulein hyperstimulation [16]. Continuing investigation will be aided greatly by employing novel analytical techniques.

Metabolomics is an exciting new analytical field in systems biology that uses sophisticated techniques such as mass spectrometry and nuclear magnetic resonance (NMR) spectroscopy to define the compliment of low molecular weight molecules present in cells, tissue or biofluids in a particular physiologic state [17]. The metabolome of a cell is the collection of downstream products of gene transcription, translation and post-translational protein modification. Thus, the metabolome may be considered a more accurate representation of the true cellular phenotype at any given time, and metabolomics represents a powerful analytical tool with which to identify cellular differences between discrete populations. NMR studies allow analysis of metabolites directly from the intact tissue. To date, no data exist regarding the pancreatic metabolome in lean or congenitally obese mice. Therefore, the aims of the current study were to describe the pancreatic lipid metabolome in lean and congenitally obese mice and to compare these results with those obtained by the conventional technique of lipid measurement using thin-layer and gas chromatography.

Materials and Methods

Animals and Diets
Twelve lean control (C57BL/6J), 12 obese leptin-deficient (Lepob) and 12 obese leptin-resistant (Lepdb) female mice were obtained from Jackson laboratory (Bar Harbor, Me., USA). At 8 weeks of age, all mice were fed a diet composed of 25% fat (soybean and corn oil), 55% carbohydrate (sucrose and cornstarch) and 20% protein (casein; Dyets Inc., Bethlehem, Pa., USA) for 4 weeks. Our laboratory has accumulated extensive baseline data under these experimental conditions. Animals were weighed weekly. All protocols for these animal studies were approved by the Indiana University Institutional Animal Care and Use Committee.

Tissue Collection
At 12 weeks of age, after an overnight fast with water allowed ad libitum, the mice were sedated with isoflurane and anesthetized with an intraperitoneal injection of xylazine (15 mg/kg) and ketamine (50 mg/kg). The animals were weighed and then underwent laparotomy and total pancreactectomy. Pancreata were frozen immediately in liquid nitrogen and preserved at −80°C for subsequent analysis by NMR (n = 6/strain) as well as by thin-layer and gas chromatography (n = 6/strain). In addition, visceral fat was harvested from the left iliopsoas fat pad of each mouse and frozen at −80°C for subsequent analysis of lipid composition by thin-layer and gas chromatography.

Metabolomic Analysis
Six pancreata from each group of mice (lean, Lepob and Lepdb) were analyzed by proton and phosphorus NMR using a CMX 400 MHz wide-bore solid spectrometer (Chemagnetics-Varian, Palo Alto, Calif., USA). The spectra of the intact tissue were obtained at 10°C by magic angle sample spinning at 3,000 Hz to remove broadening of NMR signals caused by dipolar coupling or chemical shift anisotropy. 31P NMR spectra were obtained using either a single pulse sequence without water presaturation, the 1D-NOESY sequence with water presaturation during recycle delay and mixing time, or the Carr-Purcell-Meiboom-Gill pulse sequence with water presaturation. 31P NMR spectra were obtained using a single pulse sequence. The resulting spectra were subjected to principal component analysis (PCA) to identify the metabolites with the highest variance among the different samples.

Lipid Analysis by Thin-Layer and Gas Chromatography
Pancreatic lipid content was determined by thin-layer and gas chromatography as previously described [2]. Briefly, lipids were extracted by the method of Folch-Lees. Individual lipid classes were separated by thin-layer chromatography using Silica Gel 60 A plates and visualized by rhodamine 6G. In addition, total cholesterol was analyzed by the method of Goldblatt et al. [2]. An aliquot of the Folch extract was saponified with 1 N KOH in 90% methanol. The nonsaponifiable sterol was extracted using hexane, and total cholesterol was determined using gas chromatography. The sodium salt of trimethylsilylpropionic acid, of known concentration, was used as a chemical shift as well as quantitative reference.

Lipid Quantitation by NMR
Concentrations of the total fat, triglycerides and phospholipids were determined by comparing the peak intensities of lipids,
triglycerides and choline signals, respectively, with the internal reference peak of trimethylsilylpropionic acid. The number of protons representing each peak was taken into account for the determination of the concentrations.

Statistical Analysis
Statistical analysis was performed using Sigma Stat Statistical Software (Jandel Corp., San Rafael, Calif., USA). All data are expressed as the mean ± SEM. Differences in animal body weight, pancreatic lipids and visceral lipids were tested for statistical significance by ANOVA and the Tukey test where appropriate. For analysis of the metabolome, both 1H and 31P NMR data were Fourier transformed using 4,096 data points, and the resulting spectral data were subjected to principal component multivariate statistical analysis [18] using Pirouette software version 3.11 (Infometrix Inc., Pa., USA). A p value <0.05 was considered statistically significant for all data analysis.

Results

Animal Weight
As expected, both the Lep^{ob} and Lep^{db} obese mice weighed significantly more than the lean mice (47 ± 1 and 40 ± 1 vs. 17 ± 1 g; p < 0.001). In addition, the Lep^{ob} mice weighed significantly more than the Lep^{db} mice (47 ± 1 vs. 40 ± 1; p < 0.001).

Metabolomic Analysis of Pancreatic Fat

1H NMR Data and PCA. Typical 1H NMR spectra of the pancreas of lean, Lep^{ob} and Lep^{db} mice are shown in figure 1. Dramatic biochemical differences are evident among the 3 groups. 1H NMR spectra of all tissues were dominated primarily by lipids. Specifically, much smaller intensity phosphocholine –N-CH_{3} signals (3.2 ppm) compared with lipid tail –CH_{3} clearly indicate the predominance of triglycerides over phospholipids in both obese groups of mice. Not surprisingly, the water content was significantly lower in both obese groups compared with lean mice (p < 0.001), though no differences existed in water content between the 2 groups of obese mice. As shown in figure 1b, PCA distinctly separates lean and obese (Lep^{ob} and Lep^{db}) mice based on metabolic changes. The PCA loading plot is consistent with the results that lipids and triglycerides increase in obese mice whereas phospholipids are higher in the lean mice.

31P NMR Data and PCA. 31P NMR spectra and PCA of these data are shown in figure 2. 31P NMR spectra in all tissues were dominated by resonances from inorganic phosphate (Pi), phosphomonooesters (PME) and phosphodiesters; these signals showed distinct differences among the 3 groups of pancreata. Specifically, both PME and Pi were higher in lean than in Lep^{ob} and Lep^{db} mice. The PCA analysis of these data showed a discrete clustering pattern in each strain, and the loading plot clearly indicates increased PME and Pi signals in lean mice compared with the obese groups (fig. 2b).

Total Pancreatic Fat

Chromatography. The pancreatic total fat was elevated more than 2 fold in both obese groups of mice compared with lean mice (p < 0.05). Although Lep^{db} mice had less total fat than Lep^{ob} mice, this difference was not statistically significant. Pancreatic total fat data quantitated by chromatography and metabolomics are shown in figure 3.

Metabolomics (1H NMR). Total pancreatic fat analysis by metabolomics demonstrated a pattern similar to chromatographic analysis with total fat being significantly elevated in both the obese strains of mice versus the lean mice (p < 0.05). No significant differences existed between Lep^{ob} and Lep^{db} mice in total pancreatic fat, although the pancreata of Lep^{db} mice had less total fat than those of Lep^{ob} mice.

Triglyceride Analysis

Chromatography. Pancreatic triglyceride content had a pattern similar to that of total pancreatic fat, showing a 3- to 4-fold elevation in the pancreata of both obese strains of mice versus those of lean mice (p < 0.05). No significant differences existed between Lep^{ob} and Lep^{db} mice in pancreatic triglycerides, although the Lep^{db} mice had lower triglyceride levels than the Lep^{ob} mice. Triglyceride data quantified by both chromatography and metabolomics are shown in figure 4.

Metabolomics (1H NMR). The 1H NMR analysis also showed that the pancreatic triglycerides were elevated in both obese groups of mice compared with lean animals (p < 0.05). In addition, no significant differences in pancreatic triglyceride content were detected between Lep^{ob} and Lep^{db} mice. Importantly, metabolomic analysis again corroborated the findings of conventional chromatography.

Phospholipid Analysis

Chromatography. Interestingly, pancreatic phospholipids were significantly lower in both obese strains of mice compared with the lean mice (p < 0.05). Moreover, gas chromatography showed that the pancreata of Lep^{db} mice contained significantly more phospholipids compared with Lep^{ob} mice (p < 0.05). Phospholipid analysis by both chromatography and metabolomics is shown in figure 5.
Metabolomics (1H NMR). The observation by chromatography that the pancreatic phospholipids were significantly decreased in both obese strains of mice compared with the lean mice (p < 0.05) was confirmed by metabolomic analysis. However, the chromatography finding of elevated phospholipids in Lepob mice compared with Lepob mice was not observed in the metabolomic analysis.

Free Fatty Acid Analysis

Chromatography. Total free fatty acids and individual free fatty acid chains are shown in table 1. Both saturated
Fig. 2. a $^{31}$P NMR spectra of pancreas tissue from lean, Lep$^{ob}$ and Lep$^{db}$ mice. All spectra were obtained using magic angle sample spinning NMR techniques and plotted with identical scales for direct comparison of various phosphorus metabolites in different animal models. From the comparison of peak intensities, it can be seen that PME, phosphodiester (PDE) and Pi are higher in the lean than in Lep$^{ob}$ and Lep$^{db}$. b Score plot obtained from the PCA of the $^{31}$P NMR data of Lep$^{db}$, Lep$^{ob}$ and lean mice. Both Lep$^{db}$ and Lep$^{ob}$ show distinctly separate clusters from the lean, indicating significant metabolic differences between obese and lean mice. The loading plot along the PC1 direction indicates that the phosphorus metabolites are higher in lean compared with obese mice (upward peaks).
and unsaturated free fatty acids were significantly elevated in pancreata of obese groups compared with their lean counterparts (p < 0.05). However, saturated fatty acids were significantly elevated only in Lep<sup>db</sup> mice (p < 0.05).

In addition, palmitic acid was the most abundant saturated free fatty acid chain in all 3 strains. Palmitic acid followed the same pattern as saturated free fatty acids being maximally elevated in the pancreata of Lep<sup>db</sup> mice (p < 0.05). The most abundant unsaturated free fatty acid was oleic acid which was elevated in both obese strains versus the lean mice (p < 0.05); no significant difference in oleic acid concentration existed between the obese strains.

### Table 1. Total and individual free fatty acid chains by chromatography (µg/mg pancreatic tissue)

<table>
<thead>
<tr>
<th>FFA chains</th>
<th>Lean</th>
<th>Lep&lt;sup&gt;ob&lt;/sup&gt;</th>
<th>Lep&lt;sup&gt;db&lt;/sup&gt;</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total FFA</td>
<td>2.8 ± 0.2</td>
<td>4.3 ± 0.1&lt;sup&gt;a&lt;/sup&gt;</td>
<td>5.2 ± 0.3&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Saturated FFA</td>
<td>1.2 ± 0.1</td>
<td>1.4 ± 0.2</td>
<td>1.8 ± 0.2&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Unsaturated FFA</td>
<td>1.6 ± 0.1</td>
<td>2.9 ± 0.4&lt;sup&gt;a&lt;/sup&gt;</td>
<td>3.5 ± 0.2&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Myristic (14:00)</td>
<td>0.05 ± 0.01</td>
<td>0.1 ± 0.01&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.1 ± 0.08&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Palmitic (16:00)</td>
<td>0.9 ± 0.1</td>
<td>1.1 ± 0.1</td>
<td>1.4 ± 0.1&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Palmitoleic (16:01)</td>
<td>0.2 ± 0.02</td>
<td>0.6 ± 0.07&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.7 ± 0.05&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Stearic (18:00)</td>
<td>0.3 ± 0.02</td>
<td>1.2 ± 0.03&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.3 ± 0.02&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
<tr>
<td>Oleic (18:01)</td>
<td>0.8 ± 0.08</td>
<td>1.3 ± 0.15&lt;sup&gt;a&lt;/sup&gt;</td>
<td>1.6 ± 0.1&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Linoleic (18:02)</td>
<td>0.5 ± 0.0</td>
<td>0.8 ± 0.1&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.8 ± 0.0&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Arachadonic acid</td>
<td>0.1 ± 0.01</td>
<td>0.1 ± 0.02</td>
<td>0.2 ± 0.01&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
</tbody>
</table>

*Values are the mean ± SEM. FFA = Free fatty acid.
<sup>a</sup>p < 0.05 versus lean; <sup>b</sup>p < 0.05 versus Lep<sup>ob</sup>.

Total Cholesterol and Cholesterol Ester Analysis

Chromatography. Pancreatic total cholesterol was significantly elevated in lean mice (2.2 ± 0.1 µg/mg) versus...
both Lep\textsuperscript{ob} (1.3 ± 0.1 µg/mg; p < 0.05) and Lep\textsuperscript{db} (1.8 ± 0.2 µg/mg; p < 0.05) mice. No significant difference existed between the obese strains. Additionally, no significant differences existed across the strains for cholesterol esters (lean mice = 0.6 ± 0.1, Lep\textsuperscript{ob} = 0.8 ± 0.2 and Lep\textsuperscript{db} = 1.0 ± 0.3 µg/mg; p = 0.6).

**Total Cholesterol/Phospholipid Ratio**
Across all 3 groups, no significant differences were observed in the total pancreatic cholesterol/phospholipid ratio (lean mice = 0.1 ± 0.0; Lep\textsuperscript{ob} = 0.1 ± 0.0; Lep\textsuperscript{db} = 0.1 ± 0.0; p = 0.19).

**Visceral Fat Analysis by Chromatography**
Visceral fat composition is shown in table 2.

**Visceral Triglycerides**
As expected, visceral fat composition was dominated by triglycerides. The visceral fat triglyceride content was significantly elevated in both obese strains of mice compared with their lean counterparts (p < 0.001). However, no significant differences were seen in triglyceride levels between Lep\textsuperscript{ob} and Lep\textsuperscript{db} mice. The total triglyceride content (in µg/mg tissue) was significantly greater in visceral fat compared with pancreatic fat.

**Visceral Free Fatty Acids**
Counterintuitively, total free fatty acids were decreased in the visceral fat of both obese Lep\textsuperscript{ob} (0.7 ± 0.1 µg/mg) and Lep\textsuperscript{db} (1.1 ± 0.1 µg/mg) strains compared with the lean mice (2.2 ± 0.1 µg/mg; p < 0.001). Saturated free fatty acids (lean mice = 0.7 ± 0.1, Lep\textsuperscript{ob} = 0.2 ± 0.0 and Lep\textsuperscript{db} = 0.4 ± 0.1 µg/mg) and unsaturated free fatty acids (lean mice = 1.5 ± 0.1, Lep\textsuperscript{ob} = 0.5 ± 0.1 and Lep\textsuperscript{db} = 0.7 ± 0.0 µg/mg) were similarly decreased in both obese strains of mice versus the lean mice (p < 0.001).

**Other Lipid Groups**
The preponderance of triglycerides in visceral fat samples interfered with the chromatographic analysis of the other lipid groups including phospholipids, cholesterol and cholesterol esters. Therefore, no data are available regarding the relative quantity of these lipid groups.

**Discussion**
The major finding of these experiments was that pancreata of congenitally obese mice have significantly more fat and a different composition of fat (i.e. more triglycerides and free fatty acids, and less phospholipids and cholesterol) relative to lean wild-type mice. An equally significant primary aim of this study was to validate the utility of the powerful new technique of metabolomics in this type of tissue analysis. Importantly, metabolomic analysis (by \textsuperscript{1}H and \textsuperscript{31}P NMR with PCA) correlated closely with conventional gas chromatographic analysis in identifying pancreatic total fat, triglyceride and phospholipid concentrations.

Fatty infiltration into visceral organs leads to an increased proinflammatory milieu with subsequent organ dysfunction. This process has been well documented in the cardiovascular system [19], kidney [20], gallbladder [2] and liver [1, 7]. The molecular mechanisms by which this process occurs are beginning to emerge, exemplified by the progression from steatosis to steatohepatitis to fibrosis/cirrhosis and malignancy observed in the liver [1, 7]. It is intuitive that obesity will lead to a similar situation in the pancreas; however, documentation of this phenomenon is less clear.

The association between obesity and pancreatic fat content has been shown both in human autopsy [21, 22] and recent radiology studies [23, 24]. Clinical studies have clearly shown that obesity leads to increased severity of acute pancreatitis [8–11], and obesity appears to confer an increased risk of developing pancreatic cancer [12–15]. Our group has recently shown that increased pancreatic fat is associated with an increased risk of complications after pancreatic surgery [25], and that congenitally obese (Lep\textsuperscript{ob} and Lep\textsuperscript{db}) mice develop more severe acute pancreatitis relative to lean wild-type mice [16]. Interestingly, in the latter experiments, the volume of fat per se did not correlate with severity of pancreatitis, suggesting that other factors such as alteration of the adipokine milieu or differential fat composition play a significant role in this pathological process.

<table>
<thead>
<tr>
<th></th>
<th>Lean</th>
<th>Lep\textsuperscript{ob}</th>
<th>Lep\textsuperscript{db}</th>
</tr>
</thead>
<tbody>
<tr>
<td>Triglycerides</td>
<td>459 ± 32</td>
<td>613 ± 21\textsuperscript{a}</td>
<td>663 ± 16\textsuperscript{a}</td>
</tr>
<tr>
<td>Total FFA</td>
<td>2.2 ± 0.1</td>
<td>0.7 ± 0.1\textsuperscript{a}</td>
<td>1.1 ± 0.1\textsuperscript{a}</td>
</tr>
<tr>
<td>Saturated FFA</td>
<td>0.7 ± 0.1</td>
<td>0.2 ± 0\textsuperscript{a}</td>
<td>0.4 ± 0.1\textsuperscript{a}</td>
</tr>
<tr>
<td>Unsaturated FFA</td>
<td>1.5 ± 0.1</td>
<td>0.5 ± 0.2\textsuperscript{a}</td>
<td>0.7 ± 0\textsuperscript{a}</td>
</tr>
</tbody>
</table>

Values are the mean ± SEM. FFA = Free fatty acid.
\(\textsuperscript{a}p < 0.001\) versus lean.
The current experiments showed that both congenitally obese groups of mice (Lepob and Lepdb) had significantly more total pancreatic fat, triglycerides and free fatty acids than lean wild-type mice. This finding is important, as triglycerides and free fatty acids represent the ‘toxic’ components of adipose tissue.

Triglycerides compose the largest proportion of fat in the adipocyte, and triglyceride accumulation represents the necessary ‘first hit’ towards organ dysfunction in the well-described context of nonalcoholic fatty liver disease [7]. As triglycerides accumulate, adipocyte size increases, leading to insulin resistance. Insulin resistance, in turn, promotes further triglyceride accumulation, as well as increased production of the proinflammatory cytokine TNF-α [26, 27]. Thus, this vicious cycle serves to augment the local inflammatory milieu.

Free fatty acids represent a much smaller proportion of the total lipid pool. Despite their relative paucity, free fatty acids and particularly saturated free fatty acids are critically important mediators of the proinflammatory milieu [28–30]. Circulating free fatty acids, and specifically the saturated free fatty acid palmitate, correlate positively with concentrations of the proinflammatory cytokine IL-6 [31]. In addition, animal models have shown that saturated free fatty acids preferentially upregulate the production of proinflammatory cytokines (including TNF-α and IL-6) via mechanisms mediated in part through the Toll-like receptor 4/nuclear factor-κB pathway [29, 31–35]. Furthermore, free fatty acids are an important source of reactive oxygen species through increased lipid peroxidation [36] and correlate positively with increased plasma concentrations of the proinflammatory adipokine leptin [30, 37]. Thus, our finding of increased triglycerides and free fatty acids in the pancreata of congenitally obese mice suggests that these components may play a significant role in enhancing the local inflammatory milieu.

In contrast to triglycerides and free fatty acids, both congenitally obese strains of mice had significantly less pancreatic phospholipids and cholesterol than lean animals. As both phospholipids and cholesterol are integral constituents of cell membranes, it seems likely that this observation is related to the overall increased fat volume and/or adipocyte size, with a relatively smaller proportion of cell membrane material. It was interesting to note that there were no strain differences in the total cholesterol/phospholipid ratio, suggesting that despite an overall increase in adiposity, membrane stability/fluidity remained constant.

As would be expected, visceral fat composition was dominated by triglycerides, which were significantly increased in both obese strains of mice relative to lean mice. Interestingly, the total free fatty acid volume was significantly lower in visceral fat of obese mice compared with lean animals. Further study will be necessary to see if this observation correlates with circulating free fatty acid content.

The field of metabolomics is emerging as a powerful new analytical approach in systems biology. Metabolomic analysis has been applied to toxicology studies, drug safety tests and disease detection (biomarkers). The current study represents an early effort in applying metabolomic analysis to the study of pancreatic disease, outside of 2 recent studies on pancreatic cancer [38, 39]. Our findings validate the metabolomic methodology, as patterns of total pancreatic fat, triglycerides and phospholipid concentration showed excellent correlation with those obtained by conventional thin-layer and gas chromatographic analysis. However, the real power of metabolomic analysis in evaluating pancreatic disease is yet to be recognized. Advances such as 13C NMR derivative analysis [40] promise to allow more accurate identification of metabolites unique to a specific study population (i.e. the obese mouse that develops more severe pancreatitis than a lean animal), and therefore, direct investigative focus toward specific metabolic pathways that are altered in the disease state of interest. The use of mass spectrometry especially in combination with NMR is highly promising in this direction [41–43].

Conventional chromatography methods often involve very tedious sample preparation and analysis procedures. Moreover, chromatographic methods are often selective to a certain class of metabolites, and hence, different procedures need to be employed for the analysis of a broad range of metabolites in the biological samples. On the other hand, NMR spectroscopy detects all the metabolites present in the samples, quantitatively and reproducibly. It is a high throughput analytical tool requiring less than 5 min per sample for the acquisition of the NMR data. Currently, NMR is the only technique available for profiling metabolites from intact tissue samples. Therefore, NMR spectroscopy promises immense utility in the pathophysiology of the pancreas.

In summary, these experiments demonstrate discrete differences in total fat volume and character in the pancreata of lean and congenitally obese mice. Obese mice had significantly greater pancreatic triglyceride and free fatty acid content, which may have important implications when considering the local proinflammatory milieu of the pancreas in obesity. In addition, this study highlights the utility of metabolomics-based analysis as
a powerful new analytic tool with which to define the precise mechanisms by which fat impacts pancreatic disease. Logical follow-up experiments include defining the pancreatic metabolome under various dietary conditions (i.e. low fat, high fat) as well as during pathological conditions such as pancreatitis and malignancy.

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