Loss of Fas apoptosis inhibitory molecule leads to spontaneous obesity and hepatosteatosis

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Altered hepatic lipogenesis is associated with metabolic diseases such as obesity and hepatosteatosis. Insulin resistance and compensatory hyperinsulinaemia are key drivers of these metabolic imbalances. Fas apoptosis inhibitory molecule (FAIM), a ubiquitously expressed antiapoptotic protein, functions as a mediator of Akt signalling. Since Akt acts at a nodal point in insulin signalling, we hypothesize that FAIM may be involved in energy metabolism. In the current study, C57BL/6 wild-type (WT) and FAIM-knockout (FAIM-KO) male mice were fed with normal chow diet and body weight changes were monitored. Energy expenditure, substrate utilization and physical activities were analysed using a metabolic cage. Liver, pancreas and adipose tissue were subjected to histological examination. Serum glucose and insulin levels and lipid profiles were determined by biochemical assays. Changes in components of the insulin signalling pathway in FAIM-KO mice were examined by immunoblots. We found that FAIM-KO mice developed spontaneous non-hyperphagic obesity accompanied by hepatosteatosis, adipocyte hypertrophy, dyslipidaemia, hyperglycaemia and hyperinsulinaemia. In FAIM-KO liver, lipogenesis was elevated as indicated by increased fatty acid synthesis and SREBP-1 and SREBP-2 activation. Notably, protein expression of insulin receptor beta was markedly reduced in insulin target organs of FAIM-KO mice. Akt phosphorylation was also lower in FAIM-KO liver and adipose tissue as compared with WT controls. In addition, phosphorylation of insulin receptor substrate-1 and Akt2 in response to insulin treatment in isolated FAIM-KO hepatocytes was also markedly attenuated. Altogether, our data indicate that FAIM is a novel regulator of insulin signalling and plays an essential role in energy homoeostasis. These findings may shed light on the pathogenesis of obesity and hepatosteatosis.

Cell Death and Disease (2016) 7, e2091; doi:10.1038/cddis.2016.12; published online 11 February 2016

Obesity and related metabolic diseases have reached epidemic level worldwide. Metabolic disorders including hyperglycaemia, dyslipidaemia, hypertension and central obesity often constellation as ‘metabolic syndrome’. Insulin resistance and compensatory hyperinsulinaemia underline these metabolic imbalances. Hepatic insulin resistance may stem from compromised signalling through the insulin receptor, insulin receptor substrate (IRS) and other downstream effectors such as the Akt kinase.

Liver-specific insulin receptor knockout mice exhibited insulin resistance and glucose intolerance accompanied by hyperinsulinaemia. Thus intact hepatic insulin signalling is critical for glucose metabolism. Further studies indicated that hepatic insulin resistance also accounted for dyslipidaemia and steatosis.

Hepatic insulin resistance is also associated with reduced expression of IRS-1 and IRS-2. Earlier studies showed that reduced IRS-1 expression in liver led to decreased glucokinase expression and increased blood glucose, whereas knockdown of IRS-2 resulted in elevated lipogenesis due to upregulation of SREBP-1c and fatty acid synthase and increased hepatic lipid accumulation. The dual knockdown of IRS-1 and IRS-2 resulted in systemic insulin resistance and hepatosteatosis.

As a downstream serine/threonine kinase, Akt is a critical mediator of the metabolic actions of insulin. Akt2 is the

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Abbreviations: BMI, body mass index; FAIM, Fas apoptosis inhibitory molecule; FAIM-KO, FAIM deficient; FAS, fatty acid synthase; GLUT, glucose tolerance test; HOMA-IR, homeostatic model assessment of insulin resistance; HMGCR, 3-hydroxy-3-methylglutaryl-CoA reductase; IRS, insulin receptor substrate-1; IRS-2, insulin receptor substrate-2; LDLR, low-density lipoprotein receptor; LC-MS, liquid chromatography-mass spectrometry; MRI, magnetic resonance imaging; NCD, normal chow diet; qRT-PCR, quantitative real-time PCR; SCD-1, stearoyl-CoA desaturase 1; SREBP, sterol regulatory element-binding protein; WT, wild type; XIP, X-linked inhibitor of apoptosis protein

Received 05.6.15; revised 20.11.15; accepted 08.1.16; Edited by D Vucic
predominant isoform in insulin-responsive tissues such as liver, muscle and adipose tissue. Impaired Akt signalling has been studied in livers of animal models of insulin resistance, such as ob/ob and lipodystrophic mice. Liver-specific disruption of Akt2 has mild effects on glycaemia but more dramatic effects on lipogenesis in those obese mice. Fas apoptosis inhibitory molecule (FAIM) is a critical mediator of Akt activation and was initially cloned from antigen-activated B cells and shown to protect them from activation-induced cell death. FAIM is ubiquitously expressed and conserved in evolution although it bears no homology to any known protein. Alternate splicing of FAIM generates a short (FAIM-S) and long (FAIM-L) isoform. FAIM-L is expressed in neurons. Studies demonstrate that FAIM-L contributes to inhibition of X-linked inhibitor of apoptosis protein (XIAP) ubiquitination and determines the protective or deleterious effect of tumor necrosis factor-α in neuronal cells. As a widely expressed antiapoptotic molecule, FAIM-S has been shown to participate in nuclear factor kappa-light-chain-enhancer of activated B-cell activation and prolong the lifespan of neuronal cells in culture. Our group had previously generated mice deficient in FAIM and demonstrated that FAIM played a critical role in protecting developing thymocytes and hepatocytes from Fas killing. Furthermore, we found FAIM expression to be highly expressed in symptomatic multiple myeloma patients compared with normal or premalignant individuals. Our earlier studies also showed that FAIM mediates Akt activation in thymocytes and multiple myeloma cells. Mechanistically, FAIM modulates the localization of Akt to lipid rafts during its activation. Since Akt acts as a nodal point in insulin signalling and maintains glucose and lipid homeostasis, it is conceivable that FAIM might play a role in insulin signalling and maintenance of metabolic homeostasis.

The goal of the experiments reported herein is to demonstrate that antiapoptotic molecule FAIM is a novel regulator of energy homeostasis and its expression is essential for the integrity of the insulin signalling pathway.

Results

FAIM-deficient mice develop non-hyperphagic obesity. Wild-type (WT) and FAIM-deficient (FAIM-KO) male mice were fed with normal chow diet (NCD) for 39 weeks and changes in their body weight were recorded weekly. As shown in Figure 1a, body weight gain was faster in FAIM-KO as compared with WT controls. The body weight gain in FAIM-KO mice accelerated from 19 weeks of age, and by 32 weeks, they manifested an overt obese phenotype with ~30% body weight increase compared with WT controls (Figures 1a, d and e).

To characterize energy metabolism in FAIM-KO mice, we examined their energy expenditure, substrate utilization and physical activities using metabolic cage analyses. For meaningful comparisons, WT and FAIM-KO mice subjected to metabolic cage analysis were of 14 weeks of age and had similar body weight (31.23 ± 1.72 g in WT versus...
34.06 ± 2.56 g in FAIM-KO mice, P = 0.09). FAIM-KO mice exhibited significantly reduced oxygen consumption (Figure 2a), carbon dioxide production (Figure 2b) and respiratory exchange ratio (RER) at night time (Figure 2c). No significant difference in food intake (Figure 2d) was observed, suggesting that mutant mice have spontaneous non-hyperphagic obesity. Interestingly, no reduction in loco-motor activities was observed during day and night time in mutant as compared with WT mice (Figures 2e and f). Taken together, these data suggested that reduction in resting metabolic rate, but not hyperphagia or physical inactivity, might account for the obesity seen in FAIM-deficient mice.

FAIM-deficient mice develop hypertrophic adipocytes and hepatosteatosis. To gain insight into the cause of obesity in FAIM-KO mice, we examine their fat tissue. The weight of the epididymal fat pad in FAIM-KO mice was increased ~8.4-fold (Figures 3a and b) and the size of their adipocytes was markedly enlarged (Figure 3c) as compared with WT controls. The mutant liver looked pale and slightly enlarged (Figure 3d) but there was no significant difference in the gross weight compared with WT liver. However, haematoxylin and eosin staining showed more lipid droplet accumulation in mutant livers (Figure 3f), indicating that FAIM-KO mice developed hepatic steatosis.

FAIM deficiency enhances sterol regulatory element-binding protein (SREBP) signalling and promotes lipogenesis in liver. Liquid chromatography-mass spectrometry (LC-MS)-based lipid analysis was performed to characterize fatty acid metabolism in WT and FAIM-KO livers. At 14 weeks of age, that is, before the onset of overt weight gain, saturated fatty acid [C16:0] was significantly increased in the livers of FAIM-KO mice as compared with WT controls (Supplementary Figure S1). Notably, monounsaturated fatty acids ([C16:1], [C18:1], [C20:1], [C22:1] and [C24:1]) and polyunsaturated fatty acids ([C20:2], [C18:3], [C20:3], [C18:4] and [C22:4]) were markedly elevated (Figures 4a and b), suggesting increased lipogenesis in FAIM-KO mice. Concordantly, the expression of SREBP and its downstream lipogenic target genes such as stearoyl-CoA desaturase 1 (SCD-1), fatty acid synthase (FAS) and acetyl-CoA carboxylase were all markedly increased in hepatocytes isolated from FAIM-KO mice (Figure 4c) compared with control. Quantitative real-time PCR (qRT-PCR) analysis showed that gene expressions of Srebp-1a, Srebp-1c, Acc and FAS were significantly upregulated in the mutant liver (Figure 4d). Apart from the upregulation of SREBP-1a and SREBP-1c which preferentially enhances fatty acid synthesis, we also observed upregulation of the SREBP-2 pathway that preferentially activates cholesterol synthesis in the liver by targeting 3-hydroxy-3-methyl-glutaryl-CoA reductase (HMGCR) (Figures 4e and f).

FAIM-deficient mice exhibit dyslipidaemia, hyperglycaemia and hyperinsulinaemia. We next examined the blood lipid profile of FAIM-KO mice. As compared with WT controls, total sera cholesterol, triglycerides and free fatty acids levels were all significantly elevated in the mutants from 17 weeks of age onwards (Figure 5a). Consistently, ApoB, ApoE and Ldlr mRNA expressions were also elevated in FAIM-KO livers (Figure 5b).

Since hepatosteatosis, dyslipidaemia and hyperglycaemia often cluster and give rise to ‘metabolic syndrome’, we next...
examined blood glucose levels in FAIM-KO mice. The fasting blood glucose level remained normal in FAIM-KO mice at 11 weeks of age (6.2 ± 1.3 versus 5.6 ± 0.9 mM in controls, P > 0.05). However, they exhibited fasting hyperglycaemia at 22 weeks of age (7.2 ± 1.4 versus 5.3 ± 0.5 mM in controls, P < 0.05) (Figure 5c). Further analysis by peritoneal glucose tolerance tests (GTTs) revealed that glucose disposal in skeletal muscle was normal in young (10 weeks) FAIM-KO mice whereas glucose intolerance occurred in older (50 weeks) mutants (Supplementary Figures S2a and b).

As hyperinsulinaemia underlies metabolic syndrome, we next examined fasting serum insulin levels in FAIM-KO and WT mice. Fasting serum insulin level was ~2-fold higher in mutant mice at 22 weeks of age as compared with WT controls (133.3 ± 69.0 versus 45.0 ± 17.9 pg/ml; Figure 5d). Additionally, immunohistochemical staining of insulin in pancreas revealed islet hyperplasia in FAIM-KO mice (Figure 5e). Furthermore, isolated islets from 14-week-old FAIM-KO mice displayed increased insulin secretion in response to high glucose (11 mM) stimulation in vitro (Figure 5f).

FAIM deficiency diminishes insulin actions in insulin target tissues. Hepatic insulin resistance is an important pathophysiological feature of hepatosteatosis and obesity. Hence, we attempted to elucidate the molecular mechanisms by which FAIM deficiency led to energy dysmetabolism. Insulin receptor, downstream IRS proteins and phosphatidylinositol-3-OH kinase (PI(3)K)/Akt are the main components of insulin signalling pathway. Marked reduction in insulin receptor beta (IRβ) expression was observed in ex vivo mutant liver, adipose tissues and skeletal muscle. Protein levels of IRβ, IRS-1 and IRS-2 were also significantly reduced in FAIM-KO hepatocytes as compared with WT controls (Figure 6a). Impaired Akt signalling and decreased IRS expression in the livers have been shown to be the underlying causes of insulin resistance in ob/ob and lipodystrophic mice. Our earlier studies had shown that FAIM expression was essential for the activation of Akt in thymocytes and myeloma cells. Therefore, it is conceivable that FAIM deficiency may lead to impaired Akt activation and defective insulin signalling in insulin target tissues. Indeed, Akt2 phosphorylation (Ser474) was reduced in hepatocytes and adipose tissue (Figure 6b) isolated from FAIM-KO mice. Furthermore, Akt2 (Ser474) and IRS-1 (Ser612) phosphorylation in response to insulin treatment (50 mU/ml) were also attenuated in FAIM-KO hepatocytes compared with WT controls (Figure 6c). These findings reinforced the hypothesis that FAIM plays a role in regulating insulin signalling.

FAIM expression may associate with metabolism dysregulation in human. The findings from the study of FAIM-deficient mice prompted us to examine the relevance of FAIM in human obesity and metabolic dysfunction. We analysed FAIM expression in peripheral leukocytes as they were more readily accessible than liver biopsies and had also been widely used in studies on insulin signalling and metabolic diseases. We recruited 33 obese subjects (19 females and 14 males, age 53 ± 18 years old, body mass index (BMI) 30.9 ± 4.4 kg/m², fasting blood insulin 16.2 ± 3.8 mU/l, HOMA-IR 5.1 ± 1.9) and 14 lean controls (6 females and 8 males, age 26 ± 4 years old, BMI 20.9 ± 1.7 kg/m², fasting blood insulin 9.2 ± 1.2 mU/l, HOMA-IR 2.0 ± 0.4) (Supplementary Figures S3a, c and d). FAIM expression in
leukocytes was significantly lower in obese subjects as compared with lean controls (relative expression $0.57 \pm 0.25$ versus $1.02 \pm 0.20$, $P<0.0001$) (Supplementary Figure S3b). Bivariate correlation analysis showed that FAIM expression was inversely correlated with BMI, plasma insulin levels and HOMA-IR, an established indicator of insulin resistance (Supplementary Figures S4a–c). The multivariable linear regression model further revealed that BMI and HOMA-IR were independently associated with FAIM expression after adjusting for multiple potential confounders including age (Supplementary Table S1).

**Discussion**

In our present study, we uncovered a novel role for FAIM as a metabolic regulator of insulin signalling, lipogenesis and metabolic homeostasis. We found FAIM-KO mice to develop an obese phenotype even on NCD. The spontaneous non-hyperphagic obesity in FAIM-KO mice was accompanied by enhanced lipogenesis in the liver, hypertrophic adipocytes, dyslipidaemia, hyperglycaemia and hyperinsulinaemia. The phenotype of FAIM-KO mouse resembles human metabolic syndrome which is a constellation of dyslipidaemia, hyperglycaemia, hypertension and central obesity. These observations extend our previously work by showing that, in addition to its role as a regulator of apoptosis, FAIM is critically involved in maintaining energy metabolism.

Mechanistically, we found FAIM to be involved in insulin signalling. This is evidenced by reduced phosphorylation of Akt at Ser474 in both FAIM-KO liver and adipose tissues *ex vivo* and the weakened response (resistance) to insulin treatment in FAIM-KO hepatocytes *in vitro*. These observations are consistent with our earlier studies showing that FAIM regulates Akt activation in thymocytes and myeloma cells.14,15 Therefore, the hyperinsulinaemia observed in FAIM-deficient mice might arise as compensation to insulin resistance. It also in part explains the enhanced activation of SREBP signalling and increased lipogenesis in FAIM-KO mice. Further
examination of the insulin signalling pathway in FAIM-KO mouse revealed that FAIM likely regulates the expressions of IRβ, IRS-1 and IRS-2. As shown in Figure 6a, the expressions of IRβ, IRS-1 and IRS-2 were reduced in FAIM-KO hepatocytes and IRβ protein expression was also markedly decreased in ex vivo mutant liver, adipose tissues, skeletal muscle and isolated hepatocytes. These findings extended our early finding by showing that FAIM may modulate Akt activation by regulating the expression of insulin receptor and IRS.

Interestingly, the insulin signalling defects in FAIM-KO mice were not concurrently accompanied by glucose intolerance as showed by the GTT (Supplementary Figure S2). This observation suggested that glucose disposal in skeletal muscle was not affected by the reduced insulin receptor and IRS expression in FAIM-KO mouse. This seemingly counterintuitive finding can be reconciled by earlier studies showing that glucose tolerance was normal in fat-specific and muscle-specific insulin receptor knockout mouse models. 38,39

As an evolutionally conserved and broadly expressed molecule, FAIM exists in two isoforms, with the long isoform (FAIM-L) expressed in neuronal tissues and short isoform (FAIM-S) ubiquitously expressed.21 In our mouse model, both FAIM-L and FAIM-S were deficient which prevented us from dissecting the isoform-specific function of FAIM-L and FAIM-S in the regulation of energy metabolism. This will be of interest in future studies.

Our current study may have clinical implications. Metabolic disease has become an epidemic worldwide. Insulin resistance and the associated hyperinsulinaemia are the main causes of energy dysmetabolism. Our preliminary clinical study showed that lower FAIM expression in leukocytes may be correlated with biomarkers of insulin resistance and obesity in humans. However, we cautioned that our study sample is small and there are inherent differences between lean and obese groups although we have attempted to address this issue by multivariable analysis to adjust for potential confounders such as age. On the other hand, although peripheral leukocyte has been widely used for the study of insulin signalling and metabolic disease,34–36 we are aware of the potential limitation of this methodology. Therefore, the clinical observation in our study can only be taken as hypothesis generating. A carefully planned large clinical study is

Figure 5  Dyslipidaemia, hyperglycaemia and hyperinsulinemia in FAIM-deficient mice. (a) Fasting serum concentrations of triglyceride (TG), total cholesterol (TC) and free fatty acid (FFA) in WT and FAIM−/− males (n = 3) fed with NCD at 17 weeks of age. (b) Relative mRNA expression of ApoB, ApoE and LDL receptor (Ldlr) in WT and FAIM−/− males (n = 3) fed with NCD at 17 weeks of age. Data are expressed as mean ± S.E.M. (n = 3) *P < 0.05, **P < 0.01 versus corresponding WT groups. Fasting serum glucose (c) and insulin (d) levels in WT and FAIM−/− males fed with NCD at 17 weeks of age (n = 3). (e) Representative immunohistochemical staining of insulin in the pancreas of WT and FAIM−/− males (n = 3) at 14 weeks of age. Scale bar, 100 μm. (f) Insulin production in isolated islets from WT and FAIM−/− males (n = 3) at 14 weeks of age treated with glucose (11 mM) for 2 h in culture media.
warranted to examine the role of FAIM in energy metabolism in humans.

In summary, FAIM deficiency leads to phenotypic changes in mice resembling human metabolic syndrome. Mechanistically, FAIM appears to affect insulin signalling by regulating the expression of IRβ and IRS and activation of Akt. These findings may have implications in the prevention and treatment of obesity, insulin resistance and related metabolic disorders.

Materials and Methods

Mouse maintenance and experiments. All procedures and experiments with mice were performed according to guidelines from the National Advisory Committee on Laboratory Animal Research. WT and FAIM-KO mice were housed under a 12-h light–dark cycle and given normal chow (4% of crude fat, 11% calories from fat, #1320 mod., Altromin, Large, Germany).

Mouse fat/lean composition analysis. Body composition of age-matched mutant and control littermates was measured using EchoMRI-100 (Echo Medical Systems, Houston, TX, USA).

Metabolic cages. WT and FAIM-KO male mice were housed under controlled temperature and lighting with free access to food and water. At 14 weeks of age under NCD feeding, mouse was placed individually in metabolic cages for 2 days for adaptation. Food/water intake, energy expenditure, RER and physical activity were then recorded over a period of 3 days (TSE Systems, Chesterfield, MO, USA). Data were analysed by Student’s ‘t’ test.

Mouse metabolic measurements. Mouse GTT were performed after 16 h fasting. After baseline glucose values were obtained using DiabeCHECK (Jitron, Singapore, Singapore), each mouse was given 2.0 mg glucose per gram of body weight intraperitoneally. Plasma glucose was subsequently monitored at 15, 30, 60, 90 and 120 min after the injection.

Serum glucose, free fatty acid, triglyceride, total cholesterol and insulin concentrations were measured with Glucose Colorimetric Assay Kit (Cayman Chemical), Free Fatty Acid Quantification Colorimetric/Fluorometric Kit (BioVision, Milpitas, CA, USA), Triglyceride Colorimetric Assay Kit (Cayman Chemical), Cholesterol Quantiﬁcation Kit (Sigma-Aldrich, St. Louis, MO, USA) and Ultra Sensitive Mouse Insulin ELISA Kit (CRYSTAL CHEM, Downers Grove, IL, USA), respectively.

Histology and immunohistochemistry. Mouse liver tissue, epididymal adipose and brown adipose were fixed in 10% neutral-buffered formalin, embedded in paraffin and sectioned at 5 μm. H&E staining was performed using standard techniques.

Pancreatic tissues were harvested, embedded in Tissue-Tek, and snapfrozen in dry ice and ethanol and stored at −80°C. Cryostat sections (10 μm in thickness) were prepared, air-dried and fixed in ice-cold acetone for 15 min. Sections were blocked with 5% goat serum, stained with anti-insulin antibody (Santa Cruz Biotechnology, Dallas, TX, USA), mounted with histofluid mounting medium (Paul Marienfeld GmbH & Co. KG, Lauda-Königshofen, Germany) and analysed with an

Figure 6  FAIM regulates insulin signalling and protein expressions of IRβ/IRS-1 and IRS-2. (a) Protein expression of precursor IRβ (~200 kDa) and mature IRβ (~95 kDa) in liver tissues of WT and FAIM−/− males at 14 weeks of age. IRβ expression (~95 kDa) in fat tissue and muscle of WT and FAIM−/− males at age of 14 weeks fed with NCD. N.S., nonspecific bands. Protein expression of precursor IRβ (~200 kDa), mature IRβ (~95 kDa), IRS-1 and IRS-2 in isolated hepatocytes of WT and FAIM−/− males at 14 weeks of age (lower panel). (b) Phosphorylation (Ser474) of Akt2 in isolated hepatocytes (upper) and adipose tissue (lower) of WT and FAIM−/− males fed with NCD at 14 weeks of age. (c) Insulin (50 mU/ml) induced IRS-1 (Ser612) and Akt2 (Ser474) phosphorylation in isolated WT and FAIM−/− hepatocytes (representative of three western blots).
Olympus FV1000 microscope (Olympus, Shinjuku-ku, Tokyo, Japan). Images were acquired with Olympus Fluoview Version 2.1 software.

Isolation, culture and treatment of hepatocytes. Mouse hepatocytes were isolated by two-step perfusion procedure using collagenase A according to the manufacturer’s protocol (Roche Diagnostics, Basel, Switzerland). Isolated hepatocytes were plated at 3.0×10^6 cells/ml density in gelatin-coated six-well plates (Becton Dickinson) in DMEM culture medium (1.1 g/l glucose) (Life Technologies, Rockville, MD, USA) containing MEM vitamins 1 x MEM amino acid 1 x nonessential amino acid 1 x 2 mM glutamine, 1% lactate (pH 7.4), 1% penicillin/streptomycin, 5% fetal calf serum and maintained at 37 °C under atmosphere of 5% CO₂, 95% air.

Islet isolation and treatment. Mouse pancreatic islets were extracted using a protocol modified from Bowen et al. by infusing common bile duct with collagenase P (Roche Diagnostics) and hand-picking the islets. After isolation, islets of wild-type and FAIM-deficient males were aliquoted at 10 islets per well in 48-well plates and cultured for 2 h at 37 °C in RPMI 1640 containing 10% heat-inactivated fetal calf serum, 10 mM glucose, 100 IU/ml penicillin and 100 μg/ml streptomycin. After brief centrifugation, 2 ml of the same medium was replaced. The clusters were then cultured for 2 h. Culture medium was collected and subjected to determination of insulin concentration.

Immunoblot analysis. Mouse liver, skeletal muscle or white adipose tissue clusters were then cultured for 2 h. Culture medium was collected and subjected to mouse mRNA expression analysis by qPCR. Mouse hepatocytes were plated at 3.0×10^5 cells/ml density in gelatin-coated six-well plates (Becton Dickinson) in DMEM culture medium (1.1 g/l glucose) (Life Technologies, Rockville, MD, USA) containing MEM vitamins 1 x MEM amino acid 1 x nonessential amino acid 1 x 2 mM glutamine, 1% lactate (pH 7.4), 1% penicillin/streptomycin, 5% fetal calf serum and maintained at 37 °C under atmosphere of 5% CO₂, 95% air.

LC-MS. The LC method utilized a reversed phase (C18) LC column (Acquity CSH, 2.1 x 50 mm, 1.7 μm; Waters) with two solvents: ‘A’ consisted of a mixture of acetonitrile, methanol and water (2:2:1) with 0.1% acetic acid and 0.1% ammonia solution, while ‘B’ consisted of isopropanol with 0.1% acetic acid and 0.1% ammonia solution. Samples were run according to the following LC program: the column was first equilibrated for 1 min at 1% B with a flow rate of 0.4 ml/min. The gradient was then increased from 1% B to 82.5% B over 7.5 min before B was increased to 99% for a 3.5 min wash (flow rate for wash step is 0.5 ml/min). The column was re-equilibrated for 1 min at 1% B. The column temperature was maintained at 45 °C. The eluent from the UPLC system was directed into the MS. Electrospray ionization (ESI) in the MS was conducted in both positive and negative modes in full scan with a mass range of 50–1200 m/z. The source temperature and desolvation temperature was set at 120 °C and 600 °C, respectively, while the cone gas flow and desolvation gas flow were fixed at 50 and 750 l/h, respectively. The lock mass compound was leucine enkephalin (m/z 556.2771). The capillary voltage was 2 kV for the positive ESI mode and 1 kV for the negative ESI mode.

Conflict of Interest

The authors declare no conflict of interest.

Acknowledgements

We thank Kristen Lim and Shi Ya Mak for technical assistance. We thank all volunteers and patients involved in the human FAIM expression study. This work was supported by grants from the Biomedical Research Council of the Agency for Science, Technology and Research (A-STAR), Singapore, The National Natural Science Foundation of China (No. 81373314), Research Fund for the Doctoral Program of Higher Education of China (No.20124401120012), and Alexandra Health Small Innovation Grant SIG12018C, Singapore.

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