Spexin as an indicator of beneficial effects of exercise in human obesity and diabetes

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Spexin is a novel neuropeptide playing an emerging role in metabolic diseases such as obesity and diabetes via involvement in energy homeostasis and food intake. The present study investigated the effects of obesity and type 2 diabetes (T2D) on circulating levels of spexin and its modulation by physical exercise. Normal-weight (n = 50) and obese adults with and without T2D (n = 69 and n = 66, respectively) were enrolled in the study. A subgroup of obese participants (n = 47) underwent a supervised 3-month exercise programme. Plasma spexin levels were measured by ELISA and correlated with various markers. Plasma spexin levels decreased in obese participants with or without T2D compared with those of normal-weight participants (0.43 ± 0.11, 0.44 ± 0.12 and 0.61 ± 0.23 ng/ml, respectively; \( P < 0.001 \)). Spexin levels negatively correlated with adiposity markers and blood pressure in the whole study population (\( P < 0.05 \)). Multiple regression analysis revealed blood pressure was the greatest predictive determinant of plasma spexin levels, which significantly increased in response to physical exercise in obese participants without and with T2D (\( P < 0.05 \)). Spexin levels significantly increased only in responders to exercise (those with increased oxygen consumption, \( \text{VO}_2 \text{ max} \)) with a concomitant improvement in metabolic profile. In conclusion, plasma spexin levels may be an indicator of response to physical exercise.

Energy homeostasis is regulated by complex communication pathways between the brain and peripheral organs, and disruption of these pathways significantly contributes to chronic metabolic diseases such as obesity and diabetes1. The hypothalamus produces neuropeptides that are involved in energy homeostasis. For example, genetic forms of obesity involve neuropeptides such as melanocortin-4-receptor deficiency2. Although these neuropeptides were first discovered in the brain, many are also expressed in various peripheral metabolic organs such as skeletal muscle, the pancreas and adipose tissue3. Molecules that mimic the action of neuropeptides are currently being developed for the treatment of obesity4.

Spexin, also known as neuropeptide Q, is a recently identified neuropeptide that is thought to be involved in energy homeostasis. It was primarily discovered using bioinformatics search strategies5. Studies using rodents and goldfish have shown the presence of spexin mRNA and protein in several tissues, including hypothalamus, hippocampus, cerebral cortex, stomach, kidney, adrenal gland and ovaries6. Injection of spexin into goldfish and a diet-induced obesity rodent model decreased food intake, which was linked to downregulation of orexigenic factors and upregulation of anorexigenic factors7,8. On the other hand, blood spexin levels were decreased in both adults and children with obesity and diabetes9, although another study showed no difference in serum spexin levels between three groups of adolescents (normal-weight, obese and obese with diabetes)9. Moreover, circulating spexin levels displayed a borderline association with components of metabolic syndrome in adults10. Spexin also contributes to central nervous system-mediated modulation of nociceptive responses11 and, like other neuropeptides, may also play a role in pain control. These neuropeptides were also associated with inherent differences and alterations in pain control between ethnic groups12.

Spexin gene expression was downregulated in the adipose tissue of obese patients compared with that of normal-weight controls as well as between heavy and lean co-twins13. A 1-year longitudinal study conducted in patients following Roux-en-Y gastric bypass (RYGB) in young adults with severe obesity reported increased levels of circulating spexin post-surgery14. Another nonpharmacological strategy widely used to manage obesity

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and its related complications is regular physical exercise as a component of a healthy lifestyle. Exercise has pleiotropic effects on physiology by reducing food intake and body weight as well as altering neuropeptide levels, particularly orexigenic and anorexigenic peptides, thus improving obesity-related conditions. A recent study reported favourable effects of a 6-month self-monitored lifestyle modification programme that included physical exercise on circulating spexin levels in females with prediabetes.

Taken together, these findings suggest that spexin plays a critical role in metabolic homeostasis and is potentially modulated by physical exercise. Since spexin is presumed to play a role in regulating body weight, we compared levels of circulating spexin between normal-weight, obese and obese patients with diabetes in an Arab population and evaluated the association of these levels with demographic and clinical parameters. We also investigated whether physical exercise could influence spexin levels in obese patients. Our findings indicated that levels of spexin were reduced with obesity and diabetes but increased in response to physical exercise.

**Table 1.** Anthropometric, clinical and biochemical characteristics of the study population. Data are presented as the mean ± SD. BMI, body mass index; PBF, percent body fat; SBP, systolic blood pressure; DBP, diastolic blood pressure; TC, total cholesterol; LDL, low-density lipoprotein; LDL-C, low-density lipoprotein cholesterol; TG, triglyceride; FBG, fasting blood glucose; HbA1c, hemoglobin A1c; HOMA-IR, Homeostatic Model Assessment for Insulin Resistance; hsCRP, high-sensitivity C-reactive protein.

| Characteristic           | Normal weight (N = 50) | Obese without diabetes (N = 66) | Obese with diabetes (N = 69) | P-trend | P value *versus a | P value *versus b | P value *versus c |
|--------------------------|------------------------|---------------------------------|-----------------------------|---------|------------------|------------------|------------------|
| Gender (M/F)             | 20/30                  | 27/38                           | 36/33                       | 0.304   |                  |                  |                  |
| Age (years)              | 39.96 ± 11.31          | 43.22 ± 12.44                   | 52.11 ± 9.43                | <0.001  | 0.169            | <0.001           | <0.001           |
| BMI (kg/m²)              | 23.07 ± 2.14           | 34.55 ± 3.13                    | 34.23 ± 2.56                | <0.001  | 0.001            | 0.001            | 1.000            |
| Waist (cm)               | 81.77 ± 9.86           | 107.25 ± 11.11                  | 110.19 ± 8.63               | <0.001  | <0.001           | 0.001            | 0.184            |
| Hip (cm)                 | 95.28 ± 14.64          | 116.53 ± 11.90                  | 114.60 ± 12.10              | <0.001  | <0.001           | <0.001           | 0.982            |
| PBF (%)                  | 29.08 ± 5.39           | 38.55 ± 5.51                    | 38.37 ± 8.56                | <0.001  | 0.001            | <0.001           | 1.000            |
| WBC (10⁶)                | 6.25 ± 1.46            | 6.54 ± 1.74                     | 7.43 ± 1.82                 | <0.001  | 0.832            | 0.001            | 0.001            |
| SBP (mmHg)               | 112.64 ± 10.37         | 120.21 ± 11.54                  | 122.59 ± 11.43              | <0.001  | 0.001            | <0.001           | 0.518            |
| DBP (mmHg)               | 73.94 ± 6.59           | 77.63 ± 7.64                    | 78.73 ± 6.28                | 0.001   | 0.041            | 0.001            | 0.913            |
| Resting HR (beats/min)   | 81.68 ± 11.50          | 78.88 ± 10.99                   | 82.41 ± 13.26               | 0.037   | 0.239            | 1.000            | 0.056            |
| VO₂ max (ml/kg/min)      | 21.23 ± 5.08           | 17.45 ± 4.45                    | 15.31 ± 4.07                | <0.001  | <0.001           | <0.001           | 0.007            |
| TC (mmol/l)              | 5.10 ± 0.96            | 5.16 ± 0.96                     | 4.96 ± 1.15                 | 0.365   | 1.000            | 1.000            | 0.481            |
| HDL (mmol/l)             | 1.44 ± 0.48            | 1.22 ± 0.31                     | 1.17 ± 0.39                 | <0.001  | 0.001            | <0.001           | 1.000            |
| LDL (mmol/l)             | 3.13 ± 0.85            | 3.32 ± 0.89                     | 2.99 ± 1.26                 | 0.085   | 0.751            | 1.000            | 0.082            |
| TG (mmol/l)              | 0.97 ± 0.66            | 1.37 ± 0.81                     | 1.82 ± 1.17                 | <0.001  | 0.026            | <0.001           | 0.002            |
| FBG (mmol/l)             | 5.05 ± 0.58            | 5.46 ± 0.94                     | 9.18 ± 3.57                 | <0.001  | 0.738            | <0.001           | <0.001           |
| HbA1c (%)                | 5.51 ± 0.43            | 5.77 ± 0.88                     | 8.41 ± 1.84                 | <0.001  | 0.545            | <0.001           | <0.001           |
| Insulin (ng/ml)          | 2.81 ± 1.26            | 3.89 ± 2.42                     | 4.01 ± 2.05                 | 0.007   | 0.023            | 0.010            | 1.000            |
| C-peptide (pg/ml)        | 3.86 ± 5.43            | 5.35 ± 6.41                     | 4.49 ± 5.93                 | 0.418   | 0.589            | 1.000            | 1.000            |
| HOMA-IR                  | 0.63 ± 0.28            | 0.95 ± 0.63                     | 1.50 ± 0.89                 | <0.001  | 0.063            | <0.001           | <0.001           |
| hsCRP (µg/ml)            | 3.45 ± 4.39            | 5.97 ± 4.50                     | 5.78 ± 4.35                 | 0.003   | 0.02             | 0.03             | 0.879            |

**Results**

Plasma spexin levels were dysregulated in obesity and diabetes. Anthropometric, clinical and biochemical parameters are summarised in Table 1. The obese with diabetes group was older compared with the obese without diabetes and normal-weight groups (P < 0.001). Both the obese groups (with and without diabetes) displayed significantly higher blood pressure and hsCRP levels but lower VO₂ max levels compared with the normal-weight group (P < 0.01). Moreover, the obese with diabetes group showed a more dysregulated lipid profile, as indicated by lower high-density lipoprotein (HDL) (P < 0.001) and higher TG (P < 0.001) levels, compared with the normal-weight and obese without diabetes groups. Both obese groups showed higher levels of glycermic markers (fasting blood glucose (FBG), HbA1c and insulin) (P < 0.05) compared with the normal-weight group (Table 1).

Plasma spexin levels were significantly lower in both the obese groups (P < 0.001) (Fig. 1A); however, there were no differences in levels between these two groups (P = 0.938). Not all obese individuals develop metabolic dysregulation and diabetes, and such individuals are classified as metabolically healthy individuals, despite increased body mass index (BMI). Likewise, not all non-obese individuals present a healthy metabolic profile. We divided our obese without diabetes group into metabolically healthy obese (MHO) and metabolically unhealthy obese (MUO) groups on the basis of previously reported criteria and found that levels of spexin were significantly lower in the MUO group (P = 0.043) (Fig. 1B). However, spexin levels were similar (P = 0.612) when we further divided the obese with diabetes group on the basis of HbA1c levels, controlled (HbA1c < 6.5%)
and uncontrolled (HbA1c ≥ 6.5%). The clinical and demographic characteristics of these two subcohorts are shown in Tables S1 and S2.

Spearman correlation analysis of plasma spexin levels with various demographic and clinical parameters was performed for all participants as well as separately in the normal-weight, obese without diabetes and obese with diabetes groups (Table 2). Spexin levels were inversely correlated with adiposity markers in the whole population (BMI and waist and hip circumferences) as well as with waist circumference in the normal-weight and obese with diabetes groups.

Table 2. Spearman correlation of circulating spexin with population characteristics. BMI body mass index, PBF percent body fat, SBP systolic blood pressure, DBP diastolic blood pressure, HR heart rate, VO2 max maximum oxygen consumption, TC total cholesterol, HDL high-density lipoprotein, LDL low-density lipoprotein, TG triglyceride, FBG fasting blood glucose, HbA1c hemoglobin A1c, HOMA-IR Homeostatic Model Assessment for Insulin Resistance, hsCRP high-sensitivity C-reactive protein. Significance of spearman correlation are as follows *P < 0.05, **P < 0.01.
diabetes groups. Moreover, analysis of all the study participants showed that spexin levels correlated negatively with blood pressure but positively with HDL. Within the normal-weight group, spexin levels were positively correlated with cholesterol and low-density lipoprotein (LDL) but positively correlated with HOMA-IR. On the other hand, spexin levels were positively correlated with HDL but negatively correlated with TG levels in the obese with diabetes group.

Multivariate stepwise linear regression analysis was performed with spexin as a dependent variable in the whole study population as well as separately in the normal-weight, obese without diabetes and obese with diabetes groups using Spearman correlated variables (Table 3). In the entire population and the normal-weight group, only DBP was an independent determinant of spexin levels ($P < 0.05$). However, in the obese without diabetes group, negative independent associations were observed between spexin levels and LDL and HOMA-IR ($P < 0.01$). In the obese with diabetes group, only HDL was independently associated with circulating spexin levels ($P < 0.01$).

Physical exercise increased spexin levels in the obese groups irrespective of diabetes. We assessed the effects of physical exercise by performing a pairwise comparison before and after the 3-month exercise programme of spexin levels as well as clinical and metabolic parameters on a subset of our study population [obese without diabetes ($n = 20$) and obese with diabetes ($n = 27$)]. In both obese groups, exercise significantly increased blood spexin levels ($P < 0.05$) with concurrent amelioration of clinical and metabolic markers (Fig. 2). In the obese without diabetes group, a statistically significant increase in VO2 max and decreased percent body fat and insulin levels were observed (Table 4). However, a significant improvement in heart rate (HR) with concomitant decreased levels of HbA1c and total cholesterol (TC) was observed after exercise in the obese with diabetes group (Table 5). We further performed Spearman correlation analysis for HR, VO2 max and Spexin with the other characteristics of the individuals enrolled into the exercise protocol ($n = 47$) before and after exercise (Table S3). While Spexin levels were inversely correlated with TG before exercise, they further inversely correlated with Hip circumference, insulin, HOMA-IR and hsCRP levels. As expected, HR and VO2 max inversely correlated after exercise in those individuals. Furthermore, when the two obese groups (with and without diabetes) were clustered on the basis of amelioration of cardiorespiratory fitness VO2 max after exercise (Table 6), we observed that spexin levels were only significantly increased in responders (individuals with increased VO2 max after exercise compared to their VO2 max before exercise), with decreased HR and hsCRP levels (Fig. 3).

Table 3. Multivariate linear regression analysis with spexin as dependent variable. Adjusted for age, gender and BMI.

| Independent variables | All          | Normal weight | Obese without diabetes | Obese with diabetes |
|-----------------------|--------------|---------------|------------------------|---------------------|
|                       | $\beta$ coefficient | $P$ value | $\beta$ coefficient | $P$ value |
| DBP                   | $-0.160$     | 0.040         | $-0.614$               | 0.003               |
| LDL                   | $-0.421$     | $< 0.001$     | -                      | -                   |
| HOMA-IR               | $-0.308$     | 0.009         | -                      | -                   |
| HDL                   | -            | -             | $0.375$                | 0.002               |

Figure 2. Physical exercise increased the release of spexin in obese participants. Circulating levels of Spexin were measured by ELISA using plasma samples from (A) all obese participants, or obese participants without (B) and (C) with diabetes before and after a 3-month physical exercise intervention ($n = 47$, $n = 20$ and $n = 27$, respectively). The $P$ value was determined using a paired $t$-test for intragroup comparisons before and after exercise.
Table 4. Characteristics of obese without diabetes participants before and after exercise (n = 20). Data are presented as the mean ± SD. Paired t-test was used to compare differences in participants before and after physical exercise. BMI body mass index, PBF percent body fat, SBP systolic blood pressure, DBP diastolic blood pressure, HR heart rate, VO₂ max maximum oxygen consumption, TC total cholesterol, HDL high-density lipoprotein, LDL low-density lipoprotein, TG triglyceride, FBG fasting blood glucose, HbA1c hemoglobin A1c, HOMA-IR Homeostatic Model Assessment for Insulin Resistance, hsCRP high-sensitivity C-reactive protein.

|                         | Obese without diabetes (pre) | Obese without diabetes (post) | Variation | P values |
|-------------------------|------------------------------|-------------------------------|-----------|----------|
| BMI (kg/m²)             | 34.3 ± 3.0                   | 33.9 ± 3.8                   | − 0.44 ± 0.64 | 0.196    |
| Waist (cm)              | 104.0 ± 7.6                  | 101.0 ± 6.2                  | − 2.93 ± 2.24 | 0.102    |
| Hip (cm)                | 114.8 ± 8.2                  | 111.7 ± 9.3                  | − 3.05 ± 2.17 | 0.135    |
| PBF (%)                 | 39.0 ± 4.4                   | 37.8 ± 5.5                   | − 1.21 ± 1.77 | 0.037    |
| WBC10                  | 5.9 ± 1.9                    | 5.7 ± 1.7                    | − 0.17 ± 1.18 | 0.627    |
| SBP (mmHg)              | 115.0 ± 8.2                  | 114.5 ± 6.9                  | − 0.50 ± 3.44 | 0.627    |
| DBP (mmHg)              | 75.0 ± 5.3                   | 75.0 ± 5.8                   | 0.03 ± 1.82  | 1.000    |
| Resting HR (beats/min) | 80.3 ± 8.7                   | 78.6 ± 13.6                  | − 1.70 ± 9.78 | 0.596    |
| VO₂ max (ml/kg/min)    | 17.1 ± 3.7                   | 19.8 ± 4.9                   | 2.65 ± 3.15  | 0.026    |
| TC (mmol/l)             | 5.2 ± 1.1                    | 5.1 ± 1.2                    | − 0.11 ± 0.68 | 0.499    |
| HDL (mmol/l)            | 1.3 ± 0.3                    | 1.3 ± 0.5                    | − 0.05 ± 0.26 | 0.434    |
| LDL (mmol/l)            | 3.5 ± 1.0                    | 3.2 ± 1.0                    | − 0.24 ± 0.64 | 0.127    |
| TG (mmol/l)             | 1.1 ± 0.4                    | 1.3 ± 0.9                    | 0.20 ± 0.87  | 0.159    |
| FBG (mmol/l)            | 5.3 ± 0.8                    | 5.5 ± 1.1                    | 0.16 ± 0.64  | 0.308    |
| HbA1c (%)               | 5.9 ± 1.7                    | 5.7 ± 0.5                    | − 0.20 ± 1.28 | 0.201    |
| Insulin (ng/ml)         | 4.1 ± 2.7                    | 3.2 ± 1.6                    | − 1.06 ± 2.63 | 0.039    |
| C-peptide (ng/ml)       | 8.8 ± 12.1                   | 6.3 ± 7.2                    | − 2.53 ± 3.75 | 0.138    |
| HOMA-IR                 | 1.1 ± 0.6                    | 0.8 ± 0.2                    | − 0.34 ± 0.75 | 0.070    |
| hsCRP (μg/ml)           | 5.2 ± 0.9                    | 5.7 ± 1.7                    | 0.52 ± 2.93  | 0.090    |

Table 5. Characteristics of obese with diabetes participants before and after exercise (n = 27). Data are presented as the mean ± SD. Paired t-test was used to compare differences in participants before and after physical exercise. BMI body mass index, PBF percent body fat, SBP systolic blood pressure, DBP diastolic blood pressure, HR heart rate, VO₂ max maximum oxygen consumption, TC total cholesterol, HDL high-density lipoprotein, LDL low-density lipoprotein, TG triglyceride, FBG fasting blood glucose, HbA1c hemoglobin A1c, HOMA-IR Homeostatic Model Assessment for Insulin Resistance, hsCRP high-sensitivity C-reactive protein.

|                         | Obese with diabetes (pre) | Obese with diabetes (post) | Variation | P values |
|-------------------------|----------------------------|----------------------------|-----------|----------|
| BMI (kg/m²)             | 33.8 ± 2.4                 | 33.1 ± 3.6                 | − 0.71 ± 0.59 | 0.356    |
| Waist (cm)              | 111.8 ± 5.1                | 109.5 ± 5.8                | − 2.32 ± 2.13 | 0.115    |
| Hip (cm)                | 113.5 ± 6.1                | 113.5 ± 5.6                | − 0.01 ± 0.19 | 0.988    |
| PBF (%)                 | 38.3 ± 5.1                 | 36.1 ± 5.3                 | − 2.18 ± 1.56 | 0.055    |
| WBC10                  | 7.6 ± 1.3                  | 7.4 ± 1.8                  | − 0.23 ± 0.09 | 0.622    |
| SBP (mmHg)              | 122.2 ± 13.0               | 122.6 ± 8.6                | 0.41 ± 1.38  | 0.908    |
| DBP (mmHg)              | 78.7 ± 5.0                 | 75.8 ± 5.7                 | − 0.89 ± 0.47 | 0.769    |
| Resting HR (beats/min) | 86.0 ± 7.0                 | 78.0 ± 11.7                | − 8.01 ± 6.83 | 0.042    |
| VO₂ max (ml/kg/min)    | 16.7 ± 4.2                 | 20.1 ± 4.9                 | 3.41 ± 3.89  | 0.173    |
| TC (mmol/l)             | 4.8 ± 1.1                  | 4.4 ± 0.9                  | − 0.38 ± 0.29 | 0.041    |
| HDL (mmol/l)            | 1.1 ± 0.3                  | 1.0 ± 0.3                  | − 0.07 ± 0.17 | 0.165    |
| LDL (mmol/l)            | 2.9 ± 1.1                  | 2.8 ± 0.9                  | − 0.11 ± 0.24 | 0.702    |
| TG (mmol/l)             | 1.7 ± 0.5                  | 1.6 ± 0.8                  | − 0.09 ± 0.28 | 0.799    |
| FBG (mmol/l)            | 8.6 ± 3.4                  | 8.2 ± 3.2                  | − 0.41 ± 0.88 | 0.515    |
| HbA1c (%)               | 8.2 ± 2.0                  | 7.3 ± 1.3                  | − 0.85 ± 0.63 | < 0.001  |
| Insulin (ng/ml)         | 4.0 ± 2.0                  | 3.7 ± 2.4                  | − 0.26 ± 0.49 | 0.521    |
| C-peptide (ng/ml)       | 4.4 ± 6.0                  | 4.1 ± 3.8                  | − 0.30 ± 0.14 | 0.681    |
| HOMA-IR                 | 1.6 ± 0.9                  | 1.4 ± 0.8                  | − 0.21 ± 0.37 | 0.245    |
| hsCRP (μg/ml)           | 5.3 ± 2.3                  | 6.1 ± 3.9                  | 0.81 ± 0.132  | 0.493    |
that responders displayed greater amelioration in metabolic profile. Responders showed a significant increase in fitness as seen by amelioration of VO\(_2\) max with a concomitant decrease in HR, TC, HbA1c, HOMA-IR and hsCRP. Moreover, we observed a trend of decreased BMI, waist, insulin and LDL although this did not reach statistical significance (Table 7).

### Table 6. Characteristics of all obese participants separated based on VO\(_2\) max responders after exercise. Data are presented as the mean ± SD.

| Markers        | Non-responders (N = 14) | Responders (N = 33) |
|----------------|-------------------------|---------------------|
|                | Pre                      | Post                | Variation         | P values |
| **BMI (kg/m\(^2\))** | 33.18 ± 2.17             | 32.36 ± 4.53        | −0.82 ± 1.13      | 0.189    |
| **Waist (cm)**   | 107.57 ± 6.88            | 103.14 ± 9.63       | −4.43 ± 6.58      | 0.186    |
| **Hip (cm)**     | 111.21 ± 5.79            | 108.61 ± 6.99       | −2.60 ± 1.94      | 0.304    |
| **PBF (%)**      | 36.02 ± 3.52             | 34.48 ± 4.37        | −1.54 ± 1.32      | 0.160    |
| **VO\(_2\) max** | 6.44 ± 1.32              | 5.68 ± 1.17         | −0.76 ± 0.58      | 0.139    |
| **SBP (mmHg)**  | 115.71 ± 7.87            | 116.43 ± 4.76       | 0.72 ± 1.65       | 0.805    |
| **DBP (mmHg)**  | 77.14 ± 4.88             | 77.86 ± 2.67        | 0.71 ± 1.21       | 0.788    |
| **Resting HR (beats/min)** | 83.86 ± 11.02        | 81.43 ± 14.11       | −2.43 ± 4.67      | 0.608    |
| **VO\(_2\) max (ml/kg/min)** | 17.68 ± 2.97         | 17.18 ± 3.55        | −0.50 ± 0.28      | 0.462    |
| **TC (mmol/l)** | 5.23 ± 1.14              | 5.34 ± 1.37         | 0.11 ± 0.29       | 0.767    |
| **HDL (mmol/l)**| 1.24 ± 0.34              | 1.27 ± 0.40         | 0.03 ± 0.38       | 0.649    |
| **LDL (mmol/l)**| 3.15 ± 1.30              | 3.62 ± 1.17         | 0.47 ± 0.42       | 0.384    |
| **TG (mmol/l)** | 1.35 ± 0.89              | 1.37 ± 0.77         | 0.02 ± 0.19       | 0.963    |
| **FBG (mmol/l)**| 6.37 ± 2.07              | 6.11 ± 2.13         | −0.26 ± 0.39      | 0.532    |
| **HbA1c (%)**   | 6.59 ± 1.49              | 6.34 ± 1.55         | −0.25 ± 0.27      | 0.134    |
| **Insulin (ng/ml)** | 4.81 ± 2.12            | 3.61 ± 2.11         | −1.20 ± 1.09      | 0.156    |
| **C-peptide (ng/ml)** | 11.32 ± 13.34         | 6.23 ± 7.31         | −5.09 ± 3.17      | 0.154    |
| **HOMA-IR**     | 1.72 ± 1.46              | 1.33 ± 1.15         | −0.39 ± 0.69      | 0.323    |
| **hsCRP (µg/ml)**| 2.12 ± 1.67              | 1.62 ± 1.46         | −0.50 ± 0.21      | 0.178    |

**Figure 3.** Spexin levels are increased in obese participants displaying improvement of VO\(_2\) max due to exercise. Circulating levels of Spexin were measured by ELISA using plasma samples from obese people and segregated according to increased (responders, n = 33) or not (non-responders, n = 14) VO\(_2\) max levels after 3-month physical exercise. The P value was determined using a paired t-test for intragroup comparisons before and after exercise.
In contrast, in adolescent patients with obesity or diabetes, spexin levels did not vary significantly and the obese groups and within the obese with diabetes group with controlled and uncontrolled glycemic levels normal-weight group. This was further reflected by the observed comparable levels of circulating spexin in both correlation with HOMA-IR in the obese without diabetes group and a negative correlation with HbA1c in the association between circulating spexin levels and glycemic indicators in our adult population except a positive significantly older age of the obese with diabetes group in our study, we did not observe differences in spexin levels suggests a possible role of this peptide in ageing-related functions and disorders. However, despite the significantly

A negative correlation was reported between spexin and FBG and HbA1c. Unexpectedly, we found no clear with type 2 diabetes mellitus and demonstrate that spexin could play an important role in obesity and diabetes.

Spexin, a 14-amino acid polypeptide, was first reported to suppress food intake in a goldfish model. In agreement with these findings, the present study demonstrated that circulating levels of spexin were

**Table 7.** Characteristics of all obese sorted based on spexin-responders after exercise. Data are presented as the mean ± SD. BMI, body mass index, PBF, percent body fat, SBP, systolic blood pressure, DBP, diastolic blood pressure, HOMA-IR, homeostatic model assessment of insulin resistance, hsCRP, high-sensitivity C-reactive protein.

**Discussion**

The present study showed significantly decreased plasma spexin levels in obese individuals with or without diabetes compared with healthy normal-weight individuals. Plasma spexin levels were negatively correlated with adiposity markers and blood pressure in the whole population and the normal-weight group. Furthermore, multiple regression analysis revealed that blood pressure was the most predictive determinant of plasma spexin levels among these groups. However, in the obese without diabetes group, HDL levels could predict spexin circulating levels. Interestingly, spexin levels were significantly increased by regular physical exercise in both the obese groups, and plasma spexin levels might be considered as indicator of response to physical exercise.

Spexin is a 14-amino acid polypeptide, was first reported to suppress food intake in a goldfish model. Chronic subcutaneous injection with spexin reduced food intake and led to weight loss in diet-induced obesity and brain cells. Taken together, these data highlight a key role of spexin in energy metabolism and weight regulation, with potential link to obesity and diabetes.

In agreement with these findings, the present study demonstrated that circulating levels of spexin were decreased with obesity and diabetes in adults and negatively correlated with adiposity markers (BMI and waist and hip circumference), DBP, systolic blood pressure (SBP) and lipid markers (LDL, TG, and TC) but positively correlated with HDL. These findings are in line with those of previous reports that examined the role of spexin as a metabolic regulator of body weight and glucose homeostasis in obese adults and children or patients with type 2 diabetes mellitus and demonstrate that spexin could play an important role in obesity and diabetes. A negative correlation was reported between spexin and FBG and HbA1c. Unexpectedly, we found no clear association between circulating spexin levels and glycemic indicators in our adult population except a positive correlation with HOMA-IR in the obese without diabetes group and a negative correlation with HbA1c in the normal-weight group. This was further reflected by the observed comparable levels of circulating spexin in both the obese groups and within the obese with diabetes group with controlled and uncontrolled glycemic levels (Fig. 1). In contrast, in adolescent patients with obesity or diabetes, spexin levels did not vary significantly and there was no correlation with body composition or blood measurements, indicating that spexin may not act as a metabolic regulator in adolescents.

These findings suggest that spexin might have different implications in obesity and diabetes depending on age as both of disorders are associated with age. A recent study in healthy adult women reported that circulating spexin levels were negatively correlated with age, BMI, FBG and TG. A decrease of spexin levels with age suggests a possible role of this peptide in ageing-related functions and disorders. However, despite the significantly older age of the obese with diabetes group in our study, we did not observe differences in spexin levels.
compared to the obese without diabetes group. This suggests that spexin pattern with age is blunted in diabetes status. Nevertheless, we cannot rule out the potential effects of treatment on spexin levels as study participants with diabetes were treated using antidiabetic as well as antihypertensive drugs, which might have contributed to the lack of difference of spexin between participants with and without diabetes. Very recently, Al-Daghri et al. reported increased circulating spexin levels in females with prediabetes but not in males after a 6-month self-monitored lifestyle modification programme. The same group has also reported an inverse association between spexin levels with fasting glucose in adult females with prediabetes, but not in males. This might be due to a sexual dimorphism in spexin levels in relation to gender differences in glucose metabolism and insulin sensitivity due to sex steroids. Categorisation of the obese without diabetes group into MHO and MUO groups revealed significantly lower spexin levels in the MUO group, which could not be explained by the increased glycaemic levels in the MUO. Therefore, it is crucial to elucidate the mechanisms of interaction between spexin and glucose/lipid metabolism.

Blood pressure increases with BMI. The present study showed that blood pressure (SBP and DBP) was negatively correlated with spexin and DBP regardless of BMI and was an independent determinant of spexin levels in the whole population and the normal-weight group. This may indicate the involvement of spexin in the early development of cardiovascular disease (CVD) risk in apparently healthy individuals as reflected by decreased spexin levels when CVD risk factors (BMI, blood pressure, hyperlipidemia and hyperglycemia) are increased. In line with this hypothesis, a relationship between circulating spexin levels and markers of CVDs was previously reported in adolescents with obesity, and spexin was proposed as an emerging marker for intervention and/or monitoring for CVD in children. However, no significant correlation between spexin levels and blood pressure was found in adolescents; therefore, further studies into the physiological roles and crosstalk of spexin with blood pressure are required, taking both age and BMI into consideration. It is worth noting that in our study, the negative association between spexin and DBP markers (BMI and LDL)—considered to be CVD risk factors—in the obese groups but not in the normal-weight groups endorses the protective role of spexin in metabolic disorders. In support of this hypothesis, spexin treatment reversed the dramatically increased hepatic levels of TG and total lipids in a diet-induced obesity mouse model without T2D. The positive correlation and independent association of spexin with HDL further supports the protective role of spexin as HDL decreases with obesity. Furthermore, this relationship was previously suggested in individuals with metabolic syndrome.

The present study shows for the first time that exercise increased plasma spexin levels and also improved cardiometabolic markers in obese individuals (Fig. 2). The pleiotropic beneficial effects of regular exercise are widely evidenced, especially in metabolic disorders. Indeed, regular exercise was reported in multiple studies, including ours, to improve glycaemia, insulin sensitivity and cardiometabolic markers and reduce inflammation. Interestingly, we showed that spexin levels were increased after exercise in both the obese groups with and without diabetes. We and others have previously reported greater beneficial effects of regular exercise in individuals without diabetes compared with those with diabetes based on other markers.

The inhibitory effect of spexin on body weight in diet-induced obesity mice after chronic injection of spexin was suggested to be due to a reduced respiratory exchange ratio and increased locomotor activity. Interestingly, in our study the increased levels of spexin observed after exercise were independent from BMI and diet, as only marginal changes in BMI were observed and no diet restrictions were prescribed with the exercise protocol. In line with this, spexin was reported among the BMI-independent changes observed over time in post-RYGB phase in morbidly obese youth, highlighting the fact that the clinical beneficial effects of RYGB may also involve spexin.

On the other hand, spexin and its receptor, galanin receptor 2/3 (GALR2/3), are expressed in the central nervous system as well as various peripheral tissues, such as liver, heart, adipose tissue and skeletal muscle. Since these tissues are involved in energy homeostasis and metabolic regulation, it was difficult to determine the source of increased circulating spexin after exercise in our study. Future studies are required to determine the tissue that facilitates spexin release in response to exercise to regulate body weight and food intake in obese patients with metabolic impairment.

Most previous studies using physical exercise intervention have reported the average effect of the exercise on the studied population. However, there is increased evidence that there is interindividually variability in response to exercise, as some individuals are responders who display improved metabolic markers, whereas others are non-responders and exhibit unchanged or worsened markers. Several studies have reported a high prevalence of non-responders in diabetes with or without complications, supporting a relationship between diabetes and non-response. Furthermore, MHO individuals showed higher levels of cardiorespiratory fitness than did MUO individuals, as reflected by our findings, suggesting that a healthier metabolic phenotype is linked, at least in part, to response to lifestyle factors. This variability could be also associated with other biological factors, including genetics, which account for differences in body weight regulation in response to exercise. Maximum oxygen consumption (VO₂ max or VO₂ peak) is the standard parameter of cardiorespiratory fitness and is widely used to assess the effectiveness of training and to estimate the prevalence of non-responders.

Our findings showed significantly increased spexin levels in individuals who showed improvements in VO₂ max regardless of other metabolic markers (Table 6 and Fig. 3). When we used spexin as a variable for classifying responders (with increased levels of spexin) versus non-responders, we found that more metabolic and inflammatory markers, including HR, VO₂ max, TC, HbA1c, HOMA-IR and hsCRP, were improved in the responders (Table 7). This observed parallelism between VO₂ max and spexin levels and improvement of metabolic markers in the responders highlights the potential value of spexin as a marker for exercise efficacy. Spexin, used independently or in a panel of markers, could help to tailor the intervention and thus maximise the beneficial effects of physical exercise in metabolic diseases, adapt fitness programmes for children and the elderly, or assess athletic performance. The inverse relationship between VO₂ max and HR we have observed here is in agreement with the reported link between VO₂ max and HRmax at submaximal exercise intensities.
Furthermore, the most important predictor of cardiovascular fitness and health as represented by VO2 max, was reported to be HR\textsuperscript{45}.

To the best of our knowledge, the present study is the first to report the effect of exercise on spexin levels in obesity and diabetes. This study analysed a group of high-risk obese adults with diabetes who performed supervised moderate exercise as a lifestyle approach to improve global health without diet restriction. It is worth noticing that a poor adherence to recommended diet and intensive physical activity was reported in the country\textsuperscript{46}. However, the present study had some limitations. The cross-sectional study design did not allow us to determine whether the decreased spexin levels contributed to the development of obesity and diabetes. Furthermore, the limited number of study participants did not allow generalisation of our findings. In addition, diabetes is associated with ageing, and it was challenging to find age-matched healthy normal-weight and obese controls. Moreover, our study was based on a single measurement of fasting spexin, and dietary intake was not controlled for in our study as well as the lack of detailed medication for our subjects, which may have affected spexin levels and the efficacy of physical exercise. The absence of normal-weight participants in our exercise protocol is another limitation which did not allow us to conclude on the status of spexin in those subjects. On the other hand, understanding spexin expression in adipose tissue and the effects of exercise in obese individuals with and without diabetes may elucidate the crosstalk between spexin levels and insulin resistance as well as its role in adipose tissue.

In conclusion, circulating spexin levels were decreased in human ‘diabesity’, and physical exercise restored its levels concomitant with improved cardiometabolic markers. Plasma spexin levels may be an indicator of response to physical exercise. Further studies into the impact of physical exercise in mitigating the effects of metabolic stress linked to obesity and insulin resistance on spexin are required.

**Methods**

**Study population, exercise protocol and anthropometric measurements.** The current study recruited 185 adults. Written informed consents were obtained from all participants prior to taking part in the study that was conducted in line with the principles of the Declaration of Helsinki and was approved by our Institutional Review Board. The cohort was divided into three groups: normal-weight without diabetes (n = 50), obese without diabetes (n = 66) and obese with diabetes (n = 69). As we previously reported\textsuperscript{47}, participants were excluded if they had regularly performed physical exercise within the last 6 months prior to enrolment to the study (being involved in any physical activity that is planned, structured, or repetitive at least once weekly) or had a history of major illness. Eligible participants for our 3-month supervised exercise programme were enrolled at the Fitness and Rehabilitation Centre of the Dasman Diabetes Institute\textsuperscript{47,48}.

Briefly, Prior to exercise prescription each subject underwent the Cardiopulmonary exercise test “CPET” (COSMED, Quark, Italy) using electromagnetically braked cycle ergometer to measure the maximum heart rate (max HR) and maximum oxygen consumption (VO2 max). The termination criterion of 85% of age-predicted maximal heart rate (HR max) was used to stop the test. HR max was calculated as following: HR max = 220—age (year). Physical fitness was assessed thereafter to determine the muscle strength, endurance and flexibility by performing grip strength (dynamometer), push-ups upper body strength, sit ups and forward bending test (upper and lower body flexibility). As we previously reported\textsuperscript{47}, the exercise program involved a combination of both resistance training with treadmill or cycling and moderate intensity aerobic exercise. Each exercise session included 10 min warming up and cooling down steps at 50–60% of max HR with 40 min of the mentioned program at 65–80% of max HR. Under the supervision of qualified fitness professionals at MFC, subjects exercised 3 times per week for a period of 3 months. They were monitored to maintain the recommended heart rates during the training. Anthropometric and whole-body composition measurements and exercise effectiveness was assessed at the end of the 3-month exercise program using similar fitness tests performed at baseline.

**Blood biochemical measurements.** Venous peripheral blood samples were obtained at baseline and after the 3-month exercise programme and processed as previously reported\textsuperscript{30}. Plasma biochemical and metabolic markers, including FBG and HbA1c, as well as lipid profile (TG, TC, LDL and HDL), were measured as previously reported\textsuperscript{30}. hsCRP, insulin and C-peptide levels were measured by hsCRP ELISA (Biovendor, Asheville, NC, USA), insulin and ultrasensitive C-peptide ELISA, (Mercodia AB, Uppsala, Sweden). Plasma spexin levels were measured by ELISA (#EK-023-81, Phoenix Pharmaceuticals, Burlingame, California, USA). Plasma samples were diluted 1:10 prior to spexin measurement to fit within the detection range. All measurements were performed according to the manufacturers’ instructions. Homeostatic Model Assessment of Insulin Resistance (HOMA-IR) index was calculated using the following formula: HOMA-IR = (glucose × insulin)/22.5.

**Statistical analysis.** Statistical analyses were performed using SPSS software version 25.0 (SPSS Inc., Chicago, IL, USA) as we previously reported\textsuperscript{45}. All descriptive statistics for variables in the study were reported as mean ± standard deviation. A Chi-square test was used for categorical variables, and a Wilcoxon nonparametric t-test was used for skewed variables. A two-way repeated measures ANOVA test with post hoc Bonferroni analysis was conducted to evaluate the effects of groups. A paired t-test was used to determine the significance of differences in means within each group before and after exercise. Correlations between variables were calculated using Spearman’s correlation coefficient. Multivariate linear regression analysis was performed to examine the predictive effect of each factor. Values of P < 0.05 were considered statistically significant.

**Data availability**

All generated data and resources are reported in this manuscript and there is no other data to be provided.
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Acknowledgements
We thank the staff at the Fitness and Rehabilitation Center, the Clinical Laboratory and BioBank at Dasman Diabetes Institute for their help throughout the present study. Enago (www.enago.com) is thanked for the English language review. This work was supported by the Kuwait Foundation for the Advancement of Sciences under Project (RA-2010-003).

Author contributions
A.K. and A.T. designed the study. S.K. performed the experiments. A.K., S.K. and A.T. analyzed the data, wrote and revised the manuscript. S.D. performed statistical analysis. D.M., J.A. and F.A. revised the manuscript.

Competing interests
The authors declare no competing interests.

Additional information
Supplementary information is available for this paper at https://doi.org/10.1038/s41598-020-67624-z.

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