Adiponectin in mice with altered GH action: links to insulin sensitivity and longevity?

Ellen R Lubbers¹, Edward O List¹, Adam Jara¹,², Lucila Sackman-Sala¹, Jose Cordoba-Chacon⁴, Manuel D Gahete⁵, Rhonda D Kineman⁴, Ravneet Boparai⁵, Andrzej Bartke⁵, John J Kopchick¹,² and Darlene E Berryman¹,²,³

¹Edison Biotechnology Institute, ²Department of Biomedical Sciences, College of Osteopathic Medicine and ³School of Applied Health Sciences and Wellness, College of Health Sciences and Professions, E338 Grover Center, Ohio University, Athens, Ohio 45701, USA
⁴Jesse Brown VA Medical Center, Research and Development, Chicago, Illinois 60612, USA
⁵Southern Illinois University School of Medicine, Springfield, Illinois 62794, USA

Abstract

Adiponectin is positively correlated with longevity and negatively correlated with many obesity-related diseases. While there are several circulating forms of adiponectin, the high-molecular-weight (HMW) version has been suggested to have the predominant bioactivity. Adiponectin gene expression and cognate serum protein levels are of particular interest in mice with altered GH signaling as these mice exhibit extremes in obesity that are positively associated with insulin sensitivity and lifespan as opposed to the typical negative association of these factors. While a few studies have reported total adiponectin levels in young adult mice with altered GH signaling, much remains unresolved, including changes in adiponectin levels with advancing age, proportion of total adiponectin in the HMW form, adipose depot of origin, and differential effects of GH vs IGF1. Therefore, the purpose of this study was to address these issues using assorted mouse lines with altered GH signaling. Our results show that adiponectin is generally negatively associated with GH activity, regardless of age. Further, the amount of HMW adiponectin is consistently linked with the level of total adiponectin and not necessarily with previously reported lifespan or insulin sensitivity of these mice. Interestingly, circulating adiponectin levels correlated strongly with inguinal fat mass, implying that the effects of GH on adiponectin are depot specific. Interestingly, rbGH, but not IGF1, decreased circulating total and HMW adiponectin levels. Taken together, these results fill important gaps in the literature related to GH and adiponectin and question the frequently reported associations of total and HMW adiponectin with insulin sensitivity and longevity.

Key Words
- adiponectin
- high molecular weight adiponectin
- growth hormone receptor
- growth hormone
- growth hormone deficiency
- growth hormone antagonist

Introduction

Adipose tissue, once considered a simple triglyceride storage organ, is now known as an active endocrine organ, which releases many adipokines. The most abundant adipokine synthesized and secreted from white adipose tissue (WAT) is adiponectin (Maeda et al. 1996). In humans, adiponectin has been shown to decrease in
Adiponectin in mice with altered GH action

Adiponectin is of particular interest in mice with altered GH action, as these animals exhibit alterations in obesity, insulin sensitivity, and lifespan that break the typical patterns; that is, mice with increased GH signaling tend to be lean, insulin resistant, and short-lived, while mice with low GH signaling tend to be obese, insulin sensitive, and long-lived (Coschigano et al. 2000, 2003, Berryman et al. 2004, Liu et al. 2004, Olsson et al. 2005, List et al. 2009). Previous studies have shown that adult GH receptor (Ghr<sup>−/−</sup>) knockout mice, which have essentially no GH signaling, and GH antagonist (GHA) transgenic mice, which have a reduction in GH signaling, have increased circulating total adiponectin (Berryman et al. 2004, Nilsson et al. 2005, Laron & Kopchick 2011, Masternak et al. 2012). Other mouse strains with decreased GH signaling, such as Ames dwarf, Snell dwarf, Lit/Lit, and Sma1 mice, also have elevated total adiponectin levels (Flurkey et al. 2001, Combs et al. 2003, Berryman et al. 2004, Wang et al. 2006, 2007, Arumugam et al. 2007, del Rincon et al. 2007, Alderman et al. 2009, Qiao et al. 2011).

Adiponectin is of particular interest in mice with altered GH action, as these animals exhibit alterations in obesity, insulin sensitivity, and lifespan that break the typical patterns; that is, mice with increased GH signaling tend to be lean, insulin resistant, and short-lived, while mice with low GH signaling tend to be obese, insulin sensitive, and long-lived (Coschigano et al. 2000, 2003, Berryman et al. 2004, Liu et al. 2004, Olsson et al. 2005, List et al. 2009). Previous studies have shown that adult GH receptor (Ghr<sup>−/−</sup>) knockout mice, which have essentially no GH signaling, and GH antagonist (GHA) transgenic mice, which have a reduction in GH signaling, have increased circulating total adiponectin (Berryman et al. 2004,Nilsson et al. 2005, Laron & Kopchick 2011, Masternak et al. 2012). Other mouse strains with decreased GH signaling, such as Ames dwarf, Snell dwarf, Lit/Lit, and Sma1 mice, also have elevated total adiponectin levels (Flurkey et al. 2001, Combs et al. 2003, Berryman et al. 2004, Wang et al. 2006, 2007, Arumugam et al. 2007, del Rincon et al. 2007, Alderman et al. 2009). By contrast, bGH transgenic mice have an increase in GH signaling and a decrease in circulating total adiponectin (Berryman et al. 2004, Nilsson et al. 2005, del Rincon et al. 2007, Wang et al. 2007). These mouse models of altered GH action have human clinical analogs. Ghr<sup>−/−</sup> mice are analogous to humans affected by Laron syndrome, who have a mutation in the GH receptor (Laron & Kopchick 2011). Like Ghr<sup>−/−</sup> mice, individuals with Laron syndrome have increased adiponectin (Kanety et al. 2009). Similarly, bGH transgenic mice are larger than controls with high serum levels of insulin-like growth factor 1 (IGF1) and are comparable to untreated human acromegalic individuals (Olsson et al. 2005). As in bGH mice, adiponectin levels in individuals with acromegaly are decreased (Lam et al. 2004).

There are several unresolved issues related to adiponectin in mice with altered GH action. First, <i>Ghr</i><sup>−/−</sup>, GHA, and bGH mice exhibit major changes in the amount of WAT with advancing age, which may influence adipokine secretion over time (Coschigano et al. 2000, 2003, Bartke 2003, Magon 2009, Palmer et al. 2009, Berryman et al. 2010, List et al. 2011). Secondly, little is known about the relative secretory contribution of individual WAT depots to circulating adiponectin in these mice that are known to have preferential accumulation of fat in specific depots. Thirdly, no previous studies on mice with altered GH action have differentiated between the total and high-molecular-weight (HMW) forms of adiponectin. As the HMW form is considered to have the predominant bioactivity in terms of insulin sensitivity (Pajvani et al. 2004, Fisher et al. 2005, Hara et al. 2006, Kadowaki et al. 2006, Lara-Castro et al. 2006, Trujillo & Scherer 2006, von Eynatten et al. 2008, Wang et al. 2008), studies that have assessed only total adiponectin levels may be misleading as they do not reflect the abundance of the more bioactive form. Finally, measurement of adiponectin levels in other mouse models that have more moderate alterations in the GH axis would support the link between GH action and circulating adiponectin levels. Thus, the current study includes several additional models: HiGH mice, which have a moderate (two- to threefold) increase in circulating GH; AOiGH mice, which have adult-onset isolated GH deficiency; and Ames dwarf mice, which are deficient in GH, TSH, and prolactin (Masternak et al. 2010, Gahele et al. 2011, Luque et al. 2011). Additionally, inclusion of mice injected with GH or IGF1 allows us to determine the differential effects of GH and IGF1 on circulating adiponectin. Therefore, the major goals of this study were to determine the circulating levels of total and HMW adiponectin throughout life in bGH, GHA, and <i>Ghr</i><sup>−/−</sup> mice, evaluate circulating adiponectin in other mouse lines with altered GH action, determine the depot of origin of normal and increased circulating adiponectin, examine the effects of acute GH exposure to genetically normal mice that have not been chronically exposed to altered GH levels, and establish differential effects of GH and IGF1 on circulating adiponectin.
Materials and methods

Animals and sample collection

bGH, GHA, and Ghr−/− mice For the majority of experiments, three genetically modified animal models were used: GH receptor knockout (Ghr−/−) mice, GHA mice, and bGH transgenic mice, all of which have been previously described (Chen et al. 1991, Zhou et al. 1997, Berryman et al. 2004). These three mouse strains were either produced on a pure C57BL/6j background or backcrossed more than ten generations into C57BL/6j mice. These animals were bred and housed up to four animals per cage at the Ohio University animal facility with a 10 h light:14 h darkness cycle. After weaning, mice had ad libitum access to standard rodent chow (ProLab RMH 3000, PMI Nutrition International, Inc., St Louis, MO, USA) throughout the study. Animals were fasted for 12 h overnight before whole blood collection from the tail tip using heparinized capillary tubes. Blood was centrifuged at 4000 g for 10 min at 4 °C to separate and isolate plasma, which was stored at −80 °C until time of analysis. Tissue samples were dissected after killing by cervical dislocation, flash-frozen in liquid nitrogen, and stored at −80 °C until further processing. Animal protocols for these mice were approved by Ohio University’s Institutional Animal Care and Use Committee.

Administration of GH and IGF1 C57BL/6j mice were purchased from The Jackson Laboratory at 4 weeks of age and put onto a high-fat (HF) diet (D12492; Research Diets, New Brunswick, NJ, USA). Based on previous studies, it is known that adiposity differences in chow-fed GH-injected mice are minimal, while effects of GH injection on body composition are more robust in HF-fed mice (List 2010, Ding et al. 2011). Additionally, the changes in circulating adiponectin caused by the relatively short-term exposure to GH/IGF1 in injected mice are expected to be less dramatic than the changes seen in mice with genetic modulations in the GH axis. Thus, to make potential differences in adipokine levels between groups more apparent, this study used HF-fed mice for GH and IGF1 injections. Mice were kept on a HF diet for 16 weeks and housed two to three per cage with a 10 h light:14 h darkness cycle. Purified rbGH, a gift from Monsanto (St Louis, MO, USA), and rhIGF1, a gift from Tercica, Inc. (Brisbane, CA, USA), were diluted in PBS. At 5 months of age, mice were injected s.c. twice daily with 5.0 μg rbGH/g body weight, 2.5 μg IGF1/g body weight, or both for 3 weeks. Whole blood was collected through ocular bleeding after a 12-h overnight fast. Plasma was isolated as described earlier. GH and IGF1 injection studies were conducted at Ohio University and were approved by Ohio University’s IACUC.

HiGH and AOiGHD mice Mice with elevated endogenous GH levels (referred to as HiGH mice; Gahete et al. 2011) show a selective knockout of both the Insr and IgfIr only in the GH-producing cells of the anterior pituitary, which results in a threefold increase in circulating GH levels, a 20% increase in circulating IGF1, and a modest (10%) increase in body weight compared with controls. Floxed mice served as controls. AOiGHD mice (Luque et al. 2011) have a selective destruction of the Cre-inducible diphtheria toxin receptor (iDTR) expressing GH-producing cells of the anterior pituitary following treatment with diphtheria toxin (DT) at 12 weeks of age, which leads to a 60 and 20% reduction in circulating GH and IGF1 levels respectively. DT-treated iDTR mice, not expressing the rGHPcre transgene, served as controls. Both HiGH and AOiGHD mice were created on a C57BL/6 background.

HiGH mice, AOiGHD mice, and their respective controls were housed two to four mice per cage with a 12 h light:12 h darkness cycle. Mice were weaned onto standard rodent chow (Formulab Diet, Purina Mills, Inc., Richmond, IN, USA). Based on previous studies, it is known that HiGH and AOiGHD mice show few phenotypic differences from controls when fed a low-fat diet. Thus, to make potential differences between groups more apparent, at 12 weeks of age, mice were switched to a HF diet (HF-fat, #12492; Research Diets). Ad libitum access to food and water was allowed throughout the study. At 6.5 months of age, mice were killed by decapitation and trunk blood and tissues were collected under fed conditions. The HiGH and AOiGHD studies were conducted at the Jesse Brown VA Medical Center (JBVAMC; Chicago, IL, USA) with the approval of the JBVAMC and University of Illinois at Chicago IACUC.

Ames dwarf mice Ames dwarf mice (Prop1df/Prop1df) and normal (+/+ or +/Prop1df) littermate controls were produced on a heterozygous genetic background and housed four to five per cage with a 12 h light:12 h darkness cycle. To date, there is no evidence that heterozygous animals for Ames dwarfism differ from WT animals. Mice were fed a standard chow ad libitum (Rodent Laboratory Chow 5001; LabDiet, PMI Feeds, Inc., St Louis, MO, USA). At 6 months of age, fasted whole blood was collected via cardiac puncture following isoflurane administration at Southern Illinois University. Animal protocols for these
mice were approved by the Southern Illinois University Animal Care and Use Committee.

Body composition

A quantitative NMR machine was used to analyze body composition within 1 week of blood collection in bGH, GHA, and Ghr−/− mice (Minispec, Bruker Optics, Billerica, MA, USA or Echo MRI whole-body composition analyzer; Echo Medical Systems, Houston, TX, USA).

Leptin and insulin levels

Leptin was quantified using the Quantikine Mouse Leptin Immunoassay distributed by R&D Systems (Minneapolis, MN, USA; catalog number SMOB00). Insulin was quantified using the Mouse Insulin ELISA distributed by ALPCO Diagnostics (Salem, NH, USA; catalog number 80-INSMS-E10). The intra-assay and interassay coefficients of variation (CV) were 5.3 and 7.2% respectively for insulin and 4.3 and 7.8% for leptin.

Total and HMW adiponectin in serum or plasma

HMW and total adiponectin were quantified using the Adiponectin (mouse) HMW and Total ELISA manufactured by Sekisui Medical Company (Ibaraki, Japan) and distributed by ALPCO Diagnostics (Salem, NH, USA; catalog number 47-ADPMS-E01) (Ebinuma & Matsuo 2009). Both total and HMW adiponectin are quantified on the same plate alongside a single standard curve. Total adiponectin, which includes HMW (12-mer and 18-mer), MMW (hexamer), and LMW (trimer and albumin-bound trimer), was measured without any modification to their respective structures. HMW adiponectin is measured following pretreatment of samples with a specific protease that digests hexameric and trimeric adiponectin multimers. All samples were analyzed in duplicate. Tests were performed to verify that increased lipid in the serum or plasma did not interfere with the accuracy of the assay (data not shown). The intra-assay and interassay CV were 2.7 and 3.5% respectively.

Tissue adiponectin protein content

Approximately 50 mg WAT from 6-month-old Ghr−/− and WT mice were homogenized in 150 μl PBS using a probe sonicator. Samples were centrifuged at 5000 g and the fat cake was removed from the remaining homogenate. Homogenates were stored at −80 °C until analysis with a

Table 1 Primmers for mouse Adipoq and reference genes used for quantitative real-time PCR. All primers are listed 3′–5′

<table>
<thead>
<tr>
<th>Gene</th>
<th>Forward primer</th>
<th>Reverse primer</th>
<th>Product length (bp)</th>
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<td>Adipoq</td>
<td>CTTCCTGTTCCTCTTAATCCCT</td>
<td>ACCAAGAA-GACCTGCATCTC</td>
<td>218</td>
</tr>
<tr>
<td>Rps3</td>
<td>ATCAGAGAGTT-GACCGCAGTT</td>
<td>AATGAAAGCAGAGCACACATT</td>
<td>183</td>
</tr>
<tr>
<td>B2m</td>
<td>CTGGCTTTTCTA-TATCCTGGCT</td>
<td>CATGTCTCGATCC-CAGTAGAC</td>
<td>121</td>
</tr>
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</table>

Quantikine Mouse Adiponectin/Acrp30 Immunoassay (R&D Systems; catalog number MRP300). Total protein content of the tissues was measured using the Bio-Rad Protein Assay, following the manufacturer’s specifications (Bio-Rad; catalog number 500-0006).

RNA isolation and real-time PCR

RNA was isolated from inguinal and epididymal WAT from 2-month-old bGH, 12-month-old Ghr−/−, and 18-month-old GHA mice and their WT littermate controls using TRIzol Reagent following the manufacturer’s protocol (Life Technologies; catalog number 15596-026). Sample selection was based on mouse availability and varying lifespan of these animals. cDNA was synthesized using Maxima First-Strand cDNA Synthesis Kits and quantitative real-time PCR was performed using Maxima SYBR Green/Fluorescein qPCR Master Mix (Thermo Scientific, Waltham, MA, USA). Adipoq expression in WAT was normalized to beta 2 microglobulin (B2m) and ribosomal protein S3 (Rps3). In our laboratory, these housekeeping genes have been determined to be the most stable among nine housekeeping genes for WAT (Xinyue Wang, unpublished results). The sequences of the primers used are shown in Table 1. Analysis of all qPCR data was performed with Biogazelle qbasePLUS (Biogazelle NV, Zwijnaarde, Belgium).

Statistical analysis

All data are represented as mean ± S.E.M. A multivariate two-way ANOVA with Tukey’s honestly significant difference post hoc test (SPSS 17.0, IBM SPSS Statistics, Somers, NY, USA) was used to identify differences among groups. Follow-up t-tests and one-way ANOVAs were used to determine specific genotype or tissue differences. Pearson correlations were used to analyze the relationship between insulin, leptin, tissue weights, or body composition and circulating adiponectin levels. Differences were considered significant at P<0.05.
**Results**

**Circulating HMW and total adiponectin in GHA, Ghr\(^{K/-}\), and bGH mice**

Circulating concentrations of adiponectin (HMW and total) were measured at several time points for bGH, Ghr\(^{K/-}\), and GHA mice and their controls, providing a life-long profile (Fig. 1A, B, C, D, E and F). For bGH animals, 2-, 6-, 9-, and 14-month-old mice were used; for GHA, 3.2-, 6-, 12-, and 16.5-month-old mice were used; and for Ghr\(^{K/-}\), 6-, 12-, and 24-month-old mice were used. Sample selection was based on mouse availability and varying lifespan of these animals.

The life-long total adiponectin profile shows that circulating levels of total adiponectin were increased in Ghr\(^{K/-}\) and GHA mice compared with WT controls at every time point (Fig. 1B and C). In GHA mice, total adiponectin increased over life, reaching levels comparable to Ghr\(^{K/-}\) mice by 1 year of age. Total adiponectin was significantly decreased in bGH mice compared with WT at most ages, but at 14 months of age, the difference between bGH and WT mice was no longer significant (Fig. 1A).

HMW adiponectin has not been previously reported at any age in bGH, GHA, or Ghr\(^{K/-}\) mice. Circulating total and HMW adiponectin levels were reported previously for 12- and 24-month-old WT mice (Sackmann-Sala *et al.* 2012a,b). Like total adiponectin, circulating HMW adiponectin was increased in Ghr\(^{K/-}\) mice compared with WT controls at all time points (Fig. 1F). HMW adiponectin was also increased in GHA mice compared with WT at 3.2, 12, and 16.5 months, but the increase was not significant at 6 months of age (Fig. 1E). In bGH mice, HMW adiponectin was significantly decreased in 2-month-old mice compared with WT, but there was no significant difference at older ages (Fig. 1D).

The ratio of HMW to total adiponectin has been suggested to be a valuable way to report HMW adiponectin findings, as it reflects the preferential change in HMW adiponectin production, rather than a change in all MW forms (Waki *et al.* 2003). The ratio of HMW to total adiponectin were measured side-by-side by ELISA in bGH (A and D), GHA (B and E), and Ghr\(^{K/-}\) (C and F) mice. (G, H and I) The ratio of HMW adiponectin to total adiponectin was calculated for bGH (G), GHA (H), and Ghr\(^{K/-}\) (I) mice.
Adiponectin in mice with altered GH action

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Table 2 Significant Pearson correlations in bGH, GHA, and Ghr<sup>−/−</sup> mice and their WT controls across lifespan

<table>
<thead>
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<th>Factors</th>
<th>r</th>
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<th>P</th>
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<tr>
<td>Total adiponectin × leptin</td>
<td>0.507</td>
<td>190</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Total adiponectin × insulin</td>
<td>−0.175</td>
<td>190</td>
<td>0.013</td>
</tr>
<tr>
<td>Total adiponectin × total fat</td>
<td>0.409</td>
<td>183</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Total adiponectin × inguinal</td>
<td>0.563</td>
<td>123</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>HMW adiponectin × leptin</td>
<td>0.516</td>
<td>190</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>HMW adiponectin × total fat</td>
<td>0.426</td>
<td>183</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>HMW adiponectin × inguinal</td>
<td>0.490</td>
<td>123</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>HMW/total ratio × leptin</td>
<td>0.239</td>
<td>190</td>
<td>0.001</td>
</tr>
<tr>
<td>HMW/total ratio × total fat</td>
<td>0.276</td>
<td>183</td>
<td>&lt;0.001</td>
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</table>

Adiponectin in Ghr<sup>−/−</sup> mice was significantly increased compared with WT in 6-month-old mice but was not different in 1- and 2-year-old Ghr<sup>−/−</sup> mice (Fig. 1I). GHA mice had significantly increased HMW to total adiponectin ratios at 12 and 16.5 months of age but not at 3.2 or 6 months (Fig. 1H). bGH mice and matched WT controls had similar HMW/total adiponectin ratios throughout life (Fig. 1G).

Correlational analysis

Correlational analysis was performed on all bGH, GHA, and Ghr<sup>−/−</sup> mice at all time points used in this study. Significant correlations are summarized in Table 2. The analysis revealed a strong positive (P < 0.001) correlation of total, HMW, and HMW/total adiponectin to circulating leptin concentrations. Total adiponectin was negatively correlated with circulating insulin concentrations. Correlation of total fat mass and individual depot masses with adiponectin levels revealed a strong positive (P < 0.001) correlation of total, HMW, and HMW/total adiponectin with total fat mass. Of note, total and HMW adiponectin were only significantly correlated with the mass of the inguinal fat pad (P < 0.002) but not with the mass of the epididymal or mesenteric depots.

Tissue adiponectin content

Owing to the dramatic increase in circulating adiponectin and unique WAT distribution in Ghr<sup>−/−</sup> mice with a preferential enlargement of the inguinal (subcutaneous) depot, we measured the adiponectin protein content of three adipose depots (mesenteric, inguinal, and epididymal) in 6-month-old Ghr<sup>−/−</sup> mice to determine the source of the increased circulating adiponectin (Berryman et al. 2010). Several methods of normalization of these data were considered. When normalized to total protein content, there were no differences between genotypes in adiponectin content, although there was a depot difference in Ghr<sup>−/−</sup> mice. Specifically, mesenteric WAT had significantly decreased adiponectin content (0.462 ± 0.014 ng adiponectin/μg protein) when compared with epididymal and inguinal WAT (0.879 ± 0.009 and 0.956 ± 0.130 ng adiponectin/μg protein respectively) in Ghr<sup>−/−</sup> but not in WT mice. However, normalization to total protein content does not take into account the dramatic differences in depot size and body size (therefore blood volume). To account for these differences, we normalized tissue adiponectin content to depot weight/body weight. With this method of normalization, there was a significant decrease in adiponectin in the epididymal depot and a significant increase in the inguinal depot of Ghr<sup>−/−</sup> mice when compared with WT littermate controls (Fig. 2).

Adiponectin mRNA expression

To further elucidate the potential source of changes in circulating adiponectin, adiponectin (Adipoq) expression was measured at a single time point for three models: 12-month-old Ghr<sup>−/−</sup> mice, 18-month-old GHA mice, and 2-month-old bGH mice and each group’s WT controls. It is important to note that due to the age differences between the genotypes used in this series of experiments, expression data of a given mouse strain may be compared only to its corresponding WT controls and not between transgenic lines. Adiponectin expression did not differ from WT in epididymal or inguinal WAT in 12-month-old Ghr<sup>−/−</sup>, 18-month-old GHA, or 2-month-old bGH mice (data not shown). No significant effect of
depot was found in GHA or Ghr\(^{-/-}\) mice at the ages used. However, in bGH mice but in WT mice, inguinal WAT had significantly lower adiponectin expression than epididymal WAT (Fig. 3).

**Circulating HMW and total adiponectin in Ames, HiGH, and AOiGHD mice**

To confirm the results found in bGH, GHA, and Ghr\(^{-/-}\) mice, circulating total and HMW adiponectin was also measured at about 6 months of age in three additional models of modified GH signaling (Table 3): HiGH mice, which have a more modest elevation in GH/IGF1 axis than bGH mice; AOiGHD mice, which have an adult onset GH deficiency; and Ames dwarf mice, which lack GH as well as prolactin and TSH. HiGH mice on a HF diet had a significant decrease in total and HMW adiponectin in circulation when compared with controls. However, there was no significant difference between AOiGHD mice and their controls. Ames dwarf mice showed a significant increase in total and HMW adiponectin when compared with phenotypically normal heterozygous controls. No difference in the HMW/total adiponectin ratio was observed for HiGH, AOiGHD, or Ames dwarf mice compared with their respective controls.

**Circulating HMW and total adiponectin in GH- and IGF1-injected mice**

To determine the differential effects of GH and IGF1 and to examine the effects of acute GH administration, total and HMW adiponectin was measured in HF-fed mice injected with GH and/or IGF1 (Fig. 4). Mice injected with GH or IGF1 twice daily experienced a significant decrease in total adiponectin (Fig. 4A). Mice injected with both IGF1 and GH had circulating total adiponectin levels lower than those injected with either hormone alone, indicating an additive effect of GH and IGF1 (Fig. 4A). HMW adiponectin was significantly decreased in mice injected with GH compared with controls, but was not changed in mice injected with IGF1. Mice injected with both IGF1 and GH showed a decrease in HMW adiponectin similar to those injected with only GH. Regarding the ratio of HMW to total adiponectin, mice injected with GH or GH and IGF1 had significantly decreased ratios when compared with controls (Fig. 4B).

**Discussion**

Total adiponectin levels have been previously reported to be elevated in a few strains of mice with decreased GH action, while circulating total adiponectin has been reported to decrease in mice with increased GH action (Berryman et al. 2010). Our results confirm this. All mouse models of increased GH action, bGH, HiGH, and GH-injected mice have significantly decreased total serum adiponectin levels when compared with their respective controls.
controls. By contrast, most mouse lines with decreased GH action (GHA, Ghr<sup>−/−</sup>, and Ames dwarf mice) have increased circulating total and HMW adiponectin. These findings fit nicely with data reported for individuals with acromegaly who have reduced serum adiponectin and Laron syndrome patients who have increased total and HMW adiponectin. These data highlight the importance of careful age selection when designing an experiment investigating HMW adiponectin.

No previous studies on mice with altered GH action have differentiated between total and HMW forms of adiponectin. As different forms of adiponectin may influence its bioactivity and because these mice exhibit extreme differences in longevity, insulin sensitivity, and body composition, evaluating HMW adiponectin may contribute to defining the mechanisms responsible for the differences in longevity and insulin sensitivity in these mice (Coschigano et al. 2000, 2003, Berryman et al. 2004, 2010), our data show that total adiponectin levels remain relatively constant in adult mice on a chow diet. Unlike total adiponectin, circulating levels of HMW adiponectin and the HMW/total adiponectin ratio varied with age. These data highlight the importance of careful age selection when designing an experiment investigating HMW adiponectin.

The bGH, GHA, and Ghr<sup>−/−</sup> mice have been previously shown to have significant differences in lifespan and adiposity (Coschigano et al. 2000, 2003, Bartke 2003, Magon 2009, Palmer et al. 2009, Berryman et al. 2010). Additionally, factors that affect and are affected by adipokine concentrations (e.g., adiposity and glucose tolerance) often vary with age (Arai et al. 2011, Schautz et al. 2012). Considering the variation in lifespan of these mice, comparing adipokine levels at a single time point is not appropriate, as a given chronological age may represent a different biological age in each genotype. For example, 1 year is nearing the end of a bGH mouse’s lifespan, while it is only a quarter of a Ghr<sup>−/−</sup> mouse’s lifespan and about half way through the lifespan of a WT mouse. Therefore, it might be expected that total and HMW adiponectin would vary with advancing age. Though it has been reported that adiposity increases throughout life in Ghr<sup>−/−</sup> and WT mice (Berryman et al. 2004, 2010), our data show that total adiponectin levels remain relatively constant in adult mice on a Chow diet. Unlike total adiponectin, circulating levels of HMW adiponectin and the HMW/total adiponectin ratio varied with age. These data highlight the importance of careful age selection when designing an experiment investigating HMW adiponectin.

Figure 4
Circulating total adiponectin, HMW adiponectin, and the HMW/total adiponectin ratio in HF-fed mice injected with GH, IGF1, or a combination of GH and IGF1 (Combo). Data are presented as mean ± S.E.M. Within a parameter, bars without a common letter are statistically different as determined by ANOVA. Circulating total adiponectin was measured by ELISA in mice fed a HF diet and injected with GH, IGF1, or a combination of the two (A). The ratio of HMW adiponectin to total adiponectin in mice fed a HF diet and injected with GH, IGF1, or both (B).
or longevity. These data suggest that, at least in the context of altered GH action, HMW adiponectin and HMW/total adiponectin ratio may not be strongly linked to insulin sensitivity.

As IGF1 levels are generally reflective of GH levels and many of the effects of GH act via IGF1, it is often difficult to differentiate between the effects of GH and IGF1. To shed light on this matter, we measured circulating total and HMW adiponectin in mice injected with GH, IGF1, or both. Injection with GH significantly reduced circulating total adiponectin. Interestingly, this decrease in adiponectin is associated with an improvement in many metabolic parameters, as injection of the same dose of GH has previously been shown to decrease fat mass and liver triglycerides and improve glucose tolerance (List et al. 2009). Injection with IGF1 also significantly reduced the circulating total adiponectin, though not to the extent of GH injection. Thus, we hypothesize that the actions of GH on total adiponectin levels are at least partially dependent on IGF1 action, as IGF1 injection alone lowers circulating adiponectin. Injection with both GH and IGF1 induced a greater reduction in total adiponectin than either hormone alone, implying an additive effect of GH and IGF1 on total adiponectin.

Injection with IGF1 did not change the circulating levels of HMW adiponectin or the HMW/total adiponectin ratio. These data show that while both GH and IGF1 affect total adiponectin, only GH modulates HMW adiponectin. Of note, all other mouse lines used in this study were transgenic or had natural mutations resulting in chronic alterations to GH activity. GH injections into WT mice show that an acute increase in GH action also negatively regulates circulating adiponectin levels.

The strong positive correlations between adiponectin and total fat mass suggest that these changes may be due to modulation of WAT mass by GH. As GH is lipolytic, mice with increased GH action tend to have lower WAT mass, while mice with decreased GH action have increased adipose mass (Berryman et al. 2011). Mice with low GH action show a healthy obese phenotype, escaping the negative health consequences of obesity. We hypothesize that this is through the healthy expansion of WAT without the inflammation that typically accompanies fat expansion. To support this, increased GH signaling has been associated with increased inflammation, while Ghr−/− mice have been shown to have lower levels of circulating inflammatory cytokines (Hattori 2009, Masternak et al. 2012). Likewise, inflammatory markers such as MCP1, IL6, and IL10 have been reported to increase with increased duration of HF feeding and progression of DIO (Stanton et al. 2011). We propose that the often-reported reduction of adiponectin in WT HF-fed DIO mice may be due to the inflammation associated with DIO rather than simply the increase in adipose mass. This hypothesis could also explain some of the differential effects of a HF diet on adiponectin production, as the circulating adiponectin levels would depend on both the level of inflammation and the increase in adipose mass, which could vary greatly between studies.

While it is often reported that mice with low levels of GH signaling have increased circulating adiponectin, the adipose depot responsible for the increased adiponectin is not known. As the Ghr−/− and GHA mice used in this study have a unique adipose distribution with a preferential enlargement of subcutaneous WAT and this depot is thought to have metabolically beneficial effects, it is possible that the increased adiponectin is simply due to the increased mass of the subcutaneous (inguinal) depot (Berryman et al. 2004). Additionally, the correlation of total and HMW adiponectin levels with inguinal adipose depot weights (P≤0.001) imply that this depot may be contributing more to the differences in circulating adiponectin levels. One previous study has attempted to determine the origin of elevated adiponectin in mice with decreased GH signaling. Masternak et al. (2012) performed visceral fat removal surgeries on Ghr−/− and WT mice and found that circulating adiponectin was decreased after removal of epididymal and perinephric WAT in Ghr−/− but not in WT mice, implying that visceral WAT is a primary contributor to elevated circulating adiponectin in Ghr−/− mice. This study also found that tissue adiponectin protein content normalized to total protein content is higher in epididymal WAT than in inguinal WAT. However, we found no significant genotype or depot differences in tissue adiponectin content when normalized in this manner. As mentioned previously, normalization to total protein content does not take into consideration the differences in depot and body size. In order to compensate for these differences, the current study normalized data to depot/body weight. When normalized in this manner, adiponectin content was increased in the inguinal depot of Ghr−/− mice. While these results appear to be contradictory, the Ghr−/− mice used in the visceral fat removal study have a different genetic background (129Ola/BAIb/c, C3H/C57) and, on this background, all depots, not just the inguinal fat pad, are enlarged (Panici et al. 2009). Thus, the observed difference in tissue adiponectin content could be due to different background strains. Of note, adiponectin
protein content was not assessed in all fat depots in either study. It is possible that other fat pads, such as subscapular subcutaneous fat or brown fat, could contribute substantially to the adiponectin in circulation. Thus, it is difficult to make a strong conclusion without a more comprehensive assessment. Interestingly, measurement of mRNA expression of the adiponectin gene, Adipoq, revealed no genotype or depot differences in bGH, GHA, or Ghr−/−, except that bGH mice had significantly lower expression of Adipoq in inguinal WAT when compared with epididymal. These data suggest that adiponectin is not regulated at the level of mRNA, but rather another level of regulation (translational/posttranslational/degradation) is likely responsible for the variation in circulating adiponectin levels.

While adiponectin has often been associated with increased longevity, this paper questions its importance in influencing longevity. Several models of extended longevity via alterations in the GH/IGF1 pathway have increased circulating adiponectin, including Snell dwarf, Ames dwarf, Dwarf Lit/Lit, Sma1, and Ghr−/− mice (Flurkey et al. 2001, Combs et al. 2003, Berryman et al. 2004, Wang et al. 2006, 2007, Arumugam et al. 2007, del Ronco et al. 2007, Alderman et al. 2009). Calorically restricted mice, which are also long-lived, show increased circulating adiponectin levels (McKee Alderman et al. 2010, Qiao et al. 2011). Additionally, mice expressing high levels of human adiponectin show increased longevity (Otobe et al. 2007). Links between high adiponectin levels and healthy human aging have also been reported (Arai et al. 2006, Bik et al. 2006, Atzmon et al. 2008). However, data from this study question this, as GHA mice have a normal lifespan (Coschigano et al. 2003) yet extremely high levels of circulating total and HMW adiponectin (Fig. 1B and E). These data in GHA mice demonstrate that elevated adiponectin alone is not sufficient to extend longevity in mice with reduced GH signaling. However, it should also be noted that GHA mice develop severe obesity and do not have any decrease in lifespan, indicating that the high circulating adiponectin in these mice may offer some protection from age- and obesity-associated pathologies that would normally affect obese mice.

In conclusion, the current study provides greater insight into the interplay of GH and adiponectin. The association of low GH with high adiponectin and vice versa is strengthened in this study by the measurement of adiponectin in an array of mouse strains and at varied ages. Interestingly, GH appears to modulate both total and HMW adiponectin, while IGF1 modulates only total adiponectin. The lack of consistent association of HMW adiponectin or HMW/total adiponectin ratio with insulin sensitivity sheds doubt on the relevance of these parameters as indicators of insulin sensitivity in the context of increased or decreased GH signaling. Additionally, data showing GHA mice have dramatically increased total and HMW adiponectin throughout life without extended longevity implies a need for further investigation of adiponectin’s link with aging. Using this unique system in which obesity is dissociated from insulin resistance, the current study confirms a strong negative relationship between GH signaling and adiponectin levels but questions the role of HMW adiponectin in controlling insulin sensitivity or longevity in mice.

Declaration of interest
The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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