Exploring endocrine GH pattern in mice using rank plot analysis and random blood samples

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Abstract

GH plays important pleiotropic roles in development, growth, metabolism, and aging of vertebrate species. Mouse mutants with altered GH signaling have been increasingly instrumental in studying somatotropic pathophysiology. However, the pulsatile characteristics of GH secretion are difficult to study in mice because catheterization is cumbersome and long-term serial sampling is limited by small body size and blood volume. We therefore developed an approach routinely applicable to mice, which detects endogenous, physiological GH pattern from randomly obtained spot samples. We determined individual hormone concentration in large groups of mice, ranked the data by magnitude, and statistically analyzed the resulting profiles. This revealed that the nadir-to-peak distribution of plasma GH concentration in mice was similar to other mammals, and that nycthemeral and sex differences existed as well. We found handling stress to be a potent immediate downregulator of circulating GH. We showed that samples need to be taken within seconds to reflect true endogenous levels, unaffected by stress. GH receptor/Janus kinase 2/signal transducer and activator of transcription 5 activation measured in the liver correlated strongly with plasma GH levels, but peak concentrations did not further increase the pathway activation. We applied this rank plot analysis to the GH-deficient and long-lived brain-specific IGF-1 receptor knockout (bIGF1RKO+/−) mouse mutant and found a high proportion of low GH concentrations, indicative of extended trough periods and rare peaks. Taken together, we showed that rank plot analysis is a useful method that allows straightforward studies of circadian endogenous GH levels in mice.

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Introduction

Endocrine pathophysiology and disease have been extensively modeled in spontaneous mutant and genetically engineered mice. Somatotropic hormone functions were among the first to be explored. More recently, a number of mouse mutants affecting the somatotropic axis have been used to investigate the endocrine regulation of mammalian aging and longevity, revealing that diminished GH/insulin-like growth factor 1 (IGF-1) signaling prolongs lifespan (Kenyon 2001, 2005, Bartke 2003, 2008). This has been shown using spontaneous Prop1df/df, Pit1dw/dw, and GHRHRln/ln mutants (Godfrey et al. 1993, Brown-Borg et al. 1996, Flurkey et al. 2001), GH receptor (GHR)/binding protein (BP) knockout (Coschigano et al. 2000, 2003), IGF-1R+/− and brain-specific IGF-1 receptor knockout (bIGF1RKO+/−) mutants (Holzenberger et al. 2003, Kappeler et al. 2008), and several insulin receptor substrate knockouts (Taguchi et al. 2007, Selman et al. 2008). To know more about the pleiotropic roles of endocrine GH in the pathobiology of aging, there is a strong interest to investigate GH secretion in these models. Determining the dynamic pattern of GH release in mice is, however, difficult.

Anterior pituitary somatotroph cells release GH into the blood through coordinated exocytosis following a pulsatile pattern (reviewed in Veldhuis et al. (2008) and Fig. 1A and B). Classically, the analysis of GH secretion pattern prevailing in a group of individuals requires establishing the pulsatile GH profile for each individual. This in turn requires measuring GH levels in each individual from a series of blood samples taken at short intervals (5–20 min). These samples must be taken over long periods, at least 8–10 h, to include several peaks and nadir levels, or over 24 h to allow identification of nycthemeral pattern. With this approach, the physiology of GH release has been extensively studied in humans and rats (Giustina & Veldhuis 1998). In contrast, elucidating GH pulsatile patterns in mice remains very challenging. First, because the microsurgery for venous catheterization is hazardous in small animals, in particular in growth-deficient mouse mutants, and secondly, because...
the small blood volume of mice strongly limits the number of samples that can be taken from the same individual. While larger animal species allow frequent sampling of plasma over 24 h, only one group reported on a serial sampling for GH determination in mice over 8 h (MacLeod et al. 1996). The first 600 min correspond to the dark period, and 600–1440 min to the light period. (C) Ranked distribution of the data in (A and B) confirmed low nadir and intermediate GH levels in the male compared with the female individual (P<0.005; Wilcoxon rank test). (D) On average, GH during the dark period was higher than during the light period in the female individual (P<0.005; log-rank test), whereas no difference was observed in the male.

Materials and Methods

Mice

C57Bl/6 (B6) and 129/Sv (129) mice were obtained from Charles River Laboratories (CRL, L’Arbresle, France). Igf1r<sup>floxed/+</sup> mice (Kappeler et al. 2008) were maintained in 129 genetic background for >15 generations. The Nestin-Cre (NesCre) transgene was maintained in C57Bl/6 (B6) mice for >15 generations. It produces Cre recombinase in neural and glial precursors during early neural development (Tronche et al. 1999). We mated NesCre<sup>+/−</sup> males with Igf1r<sup>floxed/floxed</sup> females to produce an F1 hybrid cohort on a 129/B6 genetic background, composed of heterozygous bIGF1RKO<sup>+/−</sup> mice and their wild-type littermate controls. Animals lived under specific pathogen free conditions in ventilated cages (Tecniplast, Milan, Italy) at 23 °C, with a 14 h light:10 h darkness cycle. Ventilated cages provide sufficient olfactory and acoustic isolation from the environment, such that the simple presence of humans in the animal room is not stressful for animals. Mice had free access to water and a commercial rodent diet. We separated mice from mothers on day 30 and housed six males or six females per cage, each cage containing three control and three mutant animals. Cages were equipped with a mouse house (Tecniplast) to enhance social interaction and prevent male aggressiveness. We conducted experiments following the institutional guidelines for the care of laboratory animals.

Genotyping

We genotyped mice by multiplex PCR using DNA from skin biopsies. Primers, 5′-CCATGGGTGTTAAATGTAA-TGGC-3′, 5′-ATGAATGCTGGTGGGTTGTTCTT-3′, and 5′-ATCTTGGAGTGGTGGGTCTGTTTC-3′, were used to amplify DNA from wild-type (256 bp), floxed (312 bp), and Cre-lox recombined (204 bp) Igf1r alleles. We detected the transgene NesCre using Cre cDNA primers 5′-CCTGGAAAATGCTTCTGTCGG-3′ and 5′-CAGGGTATTATAAGCAATCC-3′ (392 bp), and GabaR1 primers 5′-AACACACACTGCGCAGAAGCTGAGC-3′ and 5′-CAATGGTAGGCTCACTCTGGGAGATGATA-3′ (292 bp) as positive PCR control.

Blood sampling and GH RIA

GH secretion follows both nycthemeral and ultradian patterns. To detect such patterns, we collected blood samples at regular 40 min intervals over 24 h. At the time of sampling, mice were, on average, 16 weeks old, the youngest being 13 and the oldest 19 weeks old. To reduce the number of animals, but also to allow for immediate sample processing and to check for reproducibility, we performed this experiment in three sessions with a 10-day interval between them. During the first session, the mice were sampled at 0800, 1000, 1200 h, etc. the second session at 0840, 1040, 1240 h, etc. and
the third session at 0920, 1120, 1320 h, etc. At each time point, we drew blood from all six males and six females from two randomly chosen cages. Ocular sinus blood (250 μl) was collected from vigil adults using topical anesthesia and EDTA anticoagulant. For each cage, samples were taken from the first three animals immediately after opening the cages to obtain samples reflecting unstressed GH concentrations. No time was lost between moving the cage and drawing blood. The remaining three animals were sampled between 3 and 6 min after opening the cages. Using this strategy, we obtained a total of 432 samples from 144 mice (36 wild-type females, 36 bIGF1RKO+/− females, 36 wild-type males, and 36 bIGF1RKO−/− males), half of which were stressed, while the others were unstressed. Whether an individual was assigned to the stressed or unstressed group was random for each of the sessions. We thereby obtained eight groups, with mice being female or male, wild-type or mutant, and unstressed or stressed. Blood samples were immediately cooled on ice, centrifuged within 15 min, and plasma frozen and stored at −25 °C. We measured GH in 100 μl plasma per mouse using rat-specific RIA (Linco, MP Biomedical, Irvine, CA, USA) that also detects mouse GH (Kappeler et al. 2009). All samples were processed in the same assay. The sensitivity of the assay is 0.5 ng/ml. All values below this threshold were set to 0.5 ng/ml.

GHRH stimulation test

Adult males received 50 mg/kg i.p. pentobarbital anesthesia. They were injected 20 min later with mouse recombinant GHRH (0.5 mg/kg i.p., Phoenix Pharmaceuticals, Belmont, CA, USA). Blood was sampled either at the time of injection (T0) or 5 min after injection (T5), by retro-orbital puncture.

ACTH induction

ACTH was measured using Bioplex methodology (Bio-Rad). Blood samples were taken between 1000 and 1200 h to minimize any effects due to circadian rhythm. For basal ACTH, blood was sampled from unstressed mice. For stress-induced ACTH, mice were immobilized for 30 min in ventilated restraint tubes (Kent Scientific Corporation, Torrington, CT, USA) where they could not move but could breathe freely.

Tissue sampling and western blotting

To correlate the circulating GH levels with activation of the GHR/Janus kinase 2 (Jak2)/signal transducer and activator of transcription 5 (Stat5) pathway in the liver, we collected blood for RIA immediately after opening the cage, and took liver tissue samples from the same animals after exactly 2 min. Tissue samples for protein extraction were snap-frozen over liquid nitrogen. This experiment was carried out during daylight period. Western blotting was performed as described earlier (Dupont et al. 2000). We used anti-phospho-tyrosine (PY20, Transduction Laboratories, BD Biosciences, Franklin Lakes, NJ, USA), anti-mouse GHR (from G Gudmundur and F Talamanes), and anti-mouse pJak2 Tyr1007/1008 (Cell Signaling Technology, Danvers, MA, USA) antibodies. We carried out immunoprecipitation experiments with anti-rabbit Stat5 antibodies (Cell Signaling; Ozyme, Saint-Quentin-en-Yvelines, France) and immunoblotting with PY20 and then Stat5 antibodies to determine tyrosine phosphorylation of Stat5. We confirmed equal loading for each immunoblot. Bound antibody was revealed using peroxidase-conjugated secondary antibodies and ECL (Amersham Pharmacia Biotech). Signals were quantified using MacBas 2.5 (Fujifilm, Bois d’Arcy, France).

Immunohistochemistry

Pituitaries from 3-month-old bIGF1RKO+/− and control males were fixed in 4% paraformaldehyde, embedded in gelatin (Sigma), and frozen. Cryosections (18 μm) were fixed on Superfrost+ slides, and incubated with rabbit anti-GH antibody (NIDDK, National Hormone and Peptide Program), followed by incubation with Alexa-546-conjugated anti-rabbit antibody (Molecular Probes, Invitrogen, Cergy Pontoise, France). Slides were mounted with DAPI Vectashield medium (Vector Laboratories, Burlingame, CA, USA), and micrographs were acquired under identical conditions for light intensity, charge-coupled-device (CCD) image acquisition, and signal integration with a ×60 oil objective using an Olympus BX612 fluorescence microscope and DP71 CCD camera.

Statistical analysis

To rank the GH values, we assigned to each GH concentration observed in a given population the fraction of the population that has a lower GH level, similar to survival analysis. Log-rank test and the Wilcoxon signed-rank test were used to compare ranked GH distributions between the groups. The log-rank test takes account of the entire curve across all GH concentrations giving the same weight to all observations and was used to evaluate differences due to nychthemeral rhythm, stress, and Igf-1R gene mutation. The Wilcoxon signed-rank test was used for evaluating sex differences in GH because it takes into consideration preferentially the left end (nadir) of the distribution; χ² was used where indicated. The Mann–Whitney U test was used for group comparisons of the GHR/Jak/Stat pathway activation. Data showing normal distribution were analyzed by two-tailed Student’s t-test and the results were expressed as mean ± S.E.M. To estimate the number of samples needed to detect significant differences in rank plot analysis, we 1) constructed a series of pulsatile profiles and 2) applied an algorithm consisting in a random drawing of a chosen number of samples subsequently submitted to log-rank test. Differences were considered significant for P<0.05.
Results

Rank plot analysis provides a useful readout of pulsatile GH

Pincus et al. (1996) explored pulsatile GH in cannulated rats as time series (Fig. 1A and B), revealing sexual dimorphism in GH secretion. In males, large peaks occur regularly throughout the light and dark periods and are separated by long troughs. In females, large peaks occur preferentially during the dark period, with more irregular patterns of GH secretion: females produce frequent smaller peaks and higher trough values than males. We extracted the GH values from a representative male and female rat published by Pincus et al., and arranged them by order of magnitude, from lowest (nadir) to highest GH value (maximum peak; Fig. 1C). This presentation of the GH data shows the relative prevalence of low, intermediate, and high GH levels in blood, and thereby estimates the overall duration of exposure to the various GH concentrations. This ranked distribution can then be used to compare the effects of mutations or treatments on GH. In this example, the rank plot revealed low nadir levels in males ($P < 0.02$, the Wilcoxon signed-rank test). This difference depended essentially on the GH values below 15 ng/ml, i.e. nadir and intermediate GH levels. In rank plots, intermediate GH levels represent small peaks and the ascending/descending phases of high peaks. We then compared GH levels from the dark period with those from the light period. Rank plot analysis revealed lower GH levels throughout the light period in the female ($P < 0.05$, log-rank test), while there was no difference in the male (Fig. 1D). Thus, rank plot of the GH data allowed detection of known sex-related and nycthemeral differences.

We then sought to demonstrate how different GH patterns transform into rank plot. We therefore constructed six pulsatile profiles (Fig. 2A and B), representing the different types of GH secretion patterns as they can be observed due to disease, gene mutation, or aging (Pincus et al. 1996, Kappeler et al. 2004 and data herein). We then transformed these six time series into rank plots, by taking one data point every 5 min and arranging them by magnitude (Fig. 2C and D). The rank plots of these profiles were distinct from the normal pattern. Changes in the frequency of peaks or in their shape translated directly into overall elevated or suppressed GH levels, including marked changes in the proportion of nadir values. Ranked distributions from profiles with higher or lower peaks showed typical differences in the high end of GH values, not in nadir levels. Considering that a typical pulsatile profile comprises 8–10 peaks per day, about 10% of the random spot samples will fall into a peak area (compare with Fig. 1). To detect at least one peak with 95% confidence interval, the required sample size is 29. To detect the differences among various rank plot profiles with $P < 0.05$, the necessary sample size depends on the relative difference in prevalence of peak values or differences in peak values.
height. An estimation of the required sample size for a given series of differences between the groups showed that differences in peak height or frequency in the range of ± 30 to ± 40% can be revealed with sample sizes between 30 and 100 (Table 1). More subtle differences in the GH distribution require samples sizes above 100. While sample size has a clear impact on statistical power, representative rank plot profiles can be established with as few as 25 samples (Supplementary Figure 1, see section on supplementary data given at the end of this article).

Next, we used rank plot to analyze the GH data obtained in cohorts of wild-type mice. The resulting GH profiles, each representing a large group of animals, were very similar to the profiles from a single male or female rat after serial sample collection (compare Figs 3A and 1C). As in Fig. 1, the distribution in wild-type mice reflected the various phases of GH secretion, i.e. nadirs, intermediate levels, and high peaks. Most values were low, many samples were intermediate, and several high values were also present. GH values from the low range of concentrations were higher in females than in males, suggesting higher nadir levels in females. Moreover, 40% of female GH values were lower than 2.0 ng/ml, compared with 56% in males ($\chi^2=5.1, P<0.01$). As mentioned, GH secretion patterns differ during light and dark periods in many species, including rats (Pincus et al. 1996, Giustina & Veldhuis 1998). We therefore compared GH levels from the light period with those from the dark period (Fig. 3B and C).

In females, we found that GH was significantly higher during the light period ($P<0.05$; log-rank test), while differences in males were not significant. Samples were taken following a randomized 24 h schedule that allowed us to search for the nycthemeral pattern. We found no evidence for synchronization of GH secretion among individuals and no evidence for accumulation of peaks at specific times of the day.

**Table 1** Estimation of the number of observations needed for analysis of blood GH using rank plot analysis

<table>
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<tr>
<th>Quality of change</th>
<th>Change (%)</th>
<th>$n^a$</th>
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<td>Increased proportion of peaks$^b$</td>
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<td></td>
<td>-57</td>
<td>71</td>
</tr>
<tr>
<td></td>
<td>-71</td>
<td>11</td>
</tr>
<tr>
<td>Increased height of peaks$^b$</td>
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<tr>
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<td></td>
<td>+40</td>
<td>54</td>
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<td></td>
<td>+50</td>
<td>40</td>
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<tr>
<td>Decreased height of peaks$^b$</td>
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</tr>
<tr>
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<td>-30</td>
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<tr>
<td></td>
<td>-40</td>
<td>38</td>
</tr>
<tr>
<td></td>
<td>-50</td>
<td>32</td>
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$^a$Number of observations 'n' required to obtain $P<0.05$ using log-rank test.

Stress rapidly downregulates plasma GH in mice

To demonstrate the effects of physiologically stressful situations on circulating GH, we compared the GH secretion patterns from stressed and unstressed wild-type animals using rank plot analysis. Mice swiftly develop a significant stress reaction when their cage is opened and when they are handled (Balcombe et al. 2004). Therefore, in the control group, samples were taken immediately after the cage was opened, so that the GH levels still represented unstressed GH levels; in the stressed group, samples were taken 3 min later, allowing the effects of stress to develop. We found that GH values in the stressed group were predominantly in the nadir range, and that the number of intermediate values was strongly diminished compared with the unstressed (Fig. 4). In females, 75% of the GH values from stressed mice were below 2 ng/ml compared with 40% in the unstressed/control group ($\chi^2=25.1, P<0.001$); in males, 83% from stressed mice compared with 56% from the control group ($\chi^2=17.2, P<0.001$). Peak values were less frequent in stressed animals, but could rise as high as those from unstressed individuals. The overall rank plot pattern was significantly different between stressed and unstressed animals ($P<0.001$ in females, $P<0.005$ in males, log-rank test; Fig. 4).

Applying rank plot analysis to GH-deficient mouse

Next, we applied rank plot analysis to investigate GH activity in a GH-deficient mouse model. We recently reported that bFGF1RKO $^{+/-}$, using Nestin-Cre (Tronche et al. 1999)
as a Cre-lox recombination driver, selectively inhibits GH and IGF-1 signaling (Kappeler et al. 2008). These mice have low hypothalamic GHRH during development. Adult bIGF1RKO+/− have small pituitaries (males, 40% smaller than controls; females, 33%; Kappeler et al. 2008), fewer somatotroph cells (Fig. 5A), and reduced pituitary GH content (bIGF1RKO+/− males, −80%; females, −81%). To further characterize somatotroph dysfunction in this model, we measured pituitary GH release after in vivo stimulation with GHRH. This triggered significant GH secretion in the mutants and control mice, but stimulated GH levels were much lower in bIGF1RKO+/− mice than in controls (bIGF1RKO+/−, 26·4 ± 5·6 versus control, 59·7 ± 9·7 ng/ml; −44%, P < 0·01; Fig. 5B). We then investigated how the somatotrophic deficiency in bIGF1RKO+/− mice affected endogenous GH secretion. Analyzing GH distribution pattern revealed significantly lower GH levels in the mutants (P < 0·001 in both sexes; log-rank test; Fig. 5C): 75% of the GH values in bIGF1RKO+/− females were under 2·0 ng/ml versus 40% in the controls (χ² = 25·1, P < 0·001); in males, 85% of the GH values in the mutants were under 2·0 ng/ml versus 56% in the controls (χ² = 20·2, P < 0·001). Nevertheless, some very high peak values were also found in the mutants. To check whether the observed differences might be explained by any particular stress in the mutants, we measured ACTH levels, but found no change in basal hormone levels or under stress (Fig. 5D).

GHR/Jak2/Stat5 pathway activation correlates with GH levels in mice

Peripheral GHR and GHR-dependent Jak2/Stat5 signaling pathways are activated within minutes after GH peaks in the circulation (Tannenbaum et al. 2001, Verma et al. 2005, Dhir et al. 2007). We showed previously that in wild-type mice, Jak2 activation in the liver is completed 2 min after GH injection into the portal vein (Kappeler et al. 2008).

Since this delay is short compared with the time course of GH peaks, the degree of GHR/Jak2/Stat5 activation may actually correlate closely with the plasma GH concentration in mice. To investigate this, we drew blood from unstressed wild-type mice and took liver and skeletal muscle biopsies 2 min later. We then ranked these individuals by their plasma GH levels and defined four subgroups: G1–G4 (see Fig. 6A). Measuring tyrosine phosphorylation of GHR (pGHR), Jak2 (pJak2), and Stat5 (pStat5), we found a significant increase in GHR/Jak2/Stat5 pathway activation...
from nadir to low GH levels (comparing G1 with G2; Fig. 6B–E). From low to high GH levels (G2–G3), the pJak2 and pStat5 activation levels tended to increase further, but differences were not statistically significant (Fig. 6B–E). Since subgroups G1–G4 cannot be compared directly across the sexes (because of different average GH levels), we analyzed the results in a scatter plot. This revealed a strong correlation between pJak2 and plasma GH between 0.5 and 20 ng/ml for females (slope = 0.07, R = 0.73, n = 15, P < 0.001) and for males (slope = 0.11, R = 0.77, n = 15, P < 0.001; Fig. 6F). Surprisingly, GH at peak values (G4; 19.2–87.1 ng/ml) did not show higher Jak2/Stat5 pathway activation than already obtained under high GH levels (G3). We repeated this analysis in muscle, with similar result (not shown).

**Discussion**

Serial sampling by catheterization is the method of choice and gold standard for studying pulsatile GH secretion. This method, however, is difficult to implement in mouse, and only one group have reported on pulsatile GH profiles in mice (MacLeod et al. 1991, Pampori & Shapiro 1994). In this study, we have developed a method that avoids the cumbersome cannulation in small animals. We sampled blood randomly in cohorts of mice, and arranged the data as rank plots.

Comparing such data from groups of mice using log-rank and the Wilcoxon signed-rank test provided key information on GH release and somatotropic function. A powerful Approximate Entropy approach has been developed by Pincus (1991) and is used to evaluate irregularity of GH secretion (Pincus et al. 1996). However, characteristics other than irregularity are also important. In this study, we have focused on analyzing the distribution of GH concentrations, which is readily affected by mutations and treatments. Data from Low et al. (2001) and Yakar et al. (2001) on SST and LID mutant mice respectively had shown several-fold increase in average GH levels and absence of nadir values in mutants. In such cases of extreme oversecretion, GH data from spot samples can be directly compared. However, when changes are less dramatic, rank plot analysis is particularly helpful, and we propose to systematically use this approach together with log-rank or the Wilcoxon signed-rank test to evaluate GH secretion from spot samples.

To detect pulses in serial analysis in rodents, sample intervals must be 20 min or less and collected over at least 8 h to include a minimum of two peaks. Thus, the lower limit for serial GH analysis is 24 samples, per individual. In serial and random sampling, about 10% of the GH values from wild-type animals correspond to the steep parts of the peak, while the intermediate GH levels – which represent about half of the values – correspond to small peaks or

**Figure 6** Activation of the GHR/Jak2/Stat5 pathway in response to endogenous GH. (A) GH measurements were performed in blood samples from male and female wild-type mice. A strong tendency to low nadir levels in males was confirmed (P = 0.02, the Wilcoxon signed-rank test). For males and females, we then defined four subgroups according to GH levels: group 1 (G1) with nadir GH, G2 with low GH, G3 with high GH, and G4 representing peaks. (B–D) We then determined activated GHR (pGHR), pJak2, and pStat5 levels from liver samples of G1–G3 animals by western blot. Significant differences in pathway activation existed between nadir and low GH levels (G1 versus G2; n = 5 per group; *P < 0.05; **P < 0.01; NS, not significant; the Mann–Whitney U-test). (E) Representative western blots of pJak2 detection (panel C). Equivalent results were obtained for pGHR and pStat5. (F) Upper panel: scatter plot of liver pJak2 levels (G3). Lower panel: representative western blots.
the ascending/descending phases of high peaks (compare Figs 1 and 3). Based on this prevalence, statistical sample size estimation showed that about 30 determinations suffice to include elevated peak values with high probability. In practice, a representative nadir-to-peak distribution can be achieved with 30 samples, and rank plots are optimal with 50 samples. For statistical analysis, if differences in GH are several-fold and concern all parts of the ranked distribution, a nonparametric Mann–Whitney U test is sufficient. If the distribution of ranked data are to be compared, log-rank test is appropriate. If differences are mainly occurring among nadir levels, the Wilcoxon rank test should be applied to focus on comparing the nadir part of the distribution. Finally, the high 10% of the GH values can be evaluated separately by establishing linear regression. Parameters from linear regression can then be compared between mutant (or treatment) and controls, providing useful indication on prevalence, height, and broadness of peaks (not shown).

We explored the GH patterns in wild-type mice and found them similar to rats (Pincus et al. 1996, Giustina & Veldhuis 1998, Veldhuis et al. 2008). Importantly, the low nadir level – a key feature of male GH secretion – was readily picked up by rank plot analysis. This was the case when we revised the data from Pincus et al. (1996), and also when we explored ours. However, it is true that values from small and large peaks overlap in rank plot and that no clear-cut boundaries exist. Thus, rank plots allowed revealing sex differences with respect to nadir levels, but they were less sensitive concerning other known gender-related differences. Consistent with the findings from rats, we saw no significant nycthemeral differences in the GH profile in male mice. Females, in contrast, showed strong nycthemeral differences. However, GH was higher during the light period, which contrasts with the published data in rats. Interestingly, MacLeod et al. (1991) demonstrated that females have more peaks than males during the day, which is consistent with the elevated daylight GH levels of females in this study. Finally, the fact that differences between males and females in our rank plots are not as obvious as, for instance, in Fig. 1 could be due to the lower number of data points or the less sensitive GH assay. Indeed, nadir differences are equally interesting as peak values and using an assay with higher sensitivity should be considered. Moreover, it needs to be stressed that we analyzed GH in samples taken from multiple mice, thereby introducing inter-individual variation, while data in Fig. 1 compare only one male and female (see also Supplementary Figure 2, see section on supplementary data given at the end of this article).

**Rank plots reveal the effect of stress on GH release in mice**

Routine animal care, such as handling and change, causes stress and leads to marked changes in plasma hormone levels (Balcombe et al. 2004). Brown & Martin (1974) showed that in rat, plasma corticosterone increases and GH levels drop quickly after exposure to handling stress. Using rank plot analysis, we showed that mice are sensitive to handling and that cage opening was sufficient to initiate a dramatic drop in plasma GH within minutes. Even in the bIGF1RKO+/- mice, which present with GH release deficiency (Fig. 5), cage opening and handling further suppressed the plasma GH levels (not shown). This rapid GH decrease can be explained by the short initial half-life of plasma GH, which is 2-2 min in mice (Turyn & Bartke 1993) and 3-3 min in rats (Chapman et al. 1991). We conclude that wherever the mice are – in a peak or in an intermediate part of the curve – stress immediately stops secretion and GH levels fall very quickly due to the extremely short half-life of the hormone. Uptake of GH by the liver, in contrast, does not seem to play a role (Turyn et al. 1997). Interestingly, some peaks seemed resistant to stress. Consequently, to measure undisturbed GH levels, blood samples must be taken immediately upon cage opening. To further explore this swift response to stress, we reduced the time between moving the mouse and drawing blood to <20 s (experiment shown in Fig. 6A), which is roughly the time the blood needs for one circulation. Under these conditions, the overall GH distribution pattern remained the same, but the GH levels (see Fig. 6A) were on average higher than in Fig. 3, where blood was sampled within the first 2 min after cage opening. A rough estimation indicates that 40 s after starting animal handling, about 10% of the original endogenous GH is lost, and that after 1 min, this reduction may reach 20%. After 3 min, about two thirds of the original GH concentration was lost. Serial GH determination in mice by MacLeod et al. (1991) revealed a relative absence of intermediate levels, which becomes evident when data are transformed into rank plot (Supplementary Figure 2, see section on supplementary data given at the end of this article). While this particularity may be due to genetic background, it could also represent an effect of stress, similar to that reported in this study in Fig. 4. It is indeed possible that the pulses reported by MacLeod et al. correspond to the peaks that we found resistant to stress. However, these authors also showed that catheterization alone did not lead to higher stress hormone levels (MacLeod & Shapiro 1988). On the other hand, the repeated loss of blood volume itself could trigger a stress response. While it is difficult to conclude from the few existing data, these findings underscore the interest in further exploring GH pulsatility in mice. In particular, the high proportion of intermediate GH concentrations detected in spot samples from wild-type mice using rank plot analysis merits further investigation.

**Rank plot analysis of endocrine GH deficiency caused by brain IGF-1R knockout**

We reported that inactivation of IGF-1R in the embryonic brain (bIGF1RKO+/−) selectively inhibits GH and IGF-1 signaling, and that low GHRH levels are involved (Kappeler et al. 2008). The bIGF1RKO+/− mutants show depletion of pituitary GH, low circulating IGF-1, and delayed somatic growth: at 3 months of age they weigh 10% less than controls.
The bIGF1RKO \(^{+/−}\) pituitaries are smaller, display low somatotroph density, and were less responsive to GHRH stimulation. Together, these results indicated that the initial developmental defect of the hypothalamo-pituitary axis led to adult GH secretory dysfunction. Using rank plot analysis in bIGF1RKO \(^{+/−}\) mice, we showed that the hypothalamo-pituitary changes resulted in strongly reduced plasma GH. The GH pattern showed rarefied peak and more frequent trough values, suggesting that broadness and/or number of peaks were reduced in the mutants, in accordance with their diminished GHRH levels. This relationship between rank plot distribution and underlying pulsatility has been explored in Fig. 2. In bIGF1RKO \(^{+/−}\) mouse model, the marked GH deficiency in juvenile mutants is associated with reduced age-related mortality and increased lifespan. This phenotype is consistent with the findings from other long-lived mouse strains (Kenyon 2001, 2005, and references therein). Since little is known about GH release pattern in these models, we suggest exploring circulating GH in these models to gain more insight into the physiopathology of somatotropic signaling involved in mammalian longevity.

**Plasma GH and Jak/Stat pathway activation**

We investigated the activation of the GHR/Jak2/Stat5 pathway in the liver exposed to different levels of circulating GH, and found a strong correlation between circulating GH and the degree of GHR, Jak2, and Stat5 phosphorylation. This was true for GH concentrations between nadir and high GH. Similar correlation between the GH concentration and degree of Stat5 activation has been shown in rats (Choi & Waxman 2000a). The G2, G3, and G4 values (see Fig. 6) are all part of a GH pulse and it is not possible to know whether these elevated values are close to a peak or still in the ascending/descending phase of a peak. However, we demonstrated previously that Jak2 activation in wild-type mice is complete just 2 min after i.v. bolus of GH. Recently, Verma et al. (2005) showed in hypophysectomized rats that i.v. GH injection activated the Jak/Stat pathway within minutes. Importantly, we did not observe combinations of high pathway activation and low GH levels or vice versa.

Concerning GH-deficient bIGF1RKO \(^{+/−}\) mutants, we showed previously that GHR and Jak2 in the liver are hypersensitive to exogenous GH (Kappeler et al. 2008). In this study, we found that liver GHR, Jak2, and Stat5 were phosphorylated to some degree at nadir GH concentrations. Together, this could partly compensate for the reduced GH levels and explain why the growth deficit in bIGF1RKO \(^{+/−}\) mutants is relatively small compared to complete GHR knockout: adult body weight in mice with constitutive GHRKO is about 40% lower than in controls, whereas adult body weight in bIGF1RKO \(^{+/−}\) mice is only 10% less than controls (Kappeler et al. 2008, Berryman et al. 2010). Thus, it seems that even low GH concentrations together with rare peaks can support significant somatic growth.

Interestingly, the highest GH peaks that we observed (G4 in Fig. 6) did not further increase the Jak/Stat pathway activation, similar to the published data (Choi & Waxman 2000a). We suggest that the intermediate GH levels are of prime importance in the regulation of GHR pathway activation. Finally, the elevated nadir levels in females did not translate into higher Jak/Stat pathway activation, similar to rat (Choi & Waxman 1999). Similarly, male G2 and female G2 showed the same degree of activation, although GH levels in the female G2 were higher than in males (compare Fig. 6C–F). This suggests that females are somewhat resistant to endogenous GH compared to males. Others showed that Stat5b activates gender-specific gene expression in the liver in response to sex-dimorphic GH pattern (Choi & Waxman 2000b, Waxman & O’Connor 2006), a fact that is compatible with our data. Studying these mechanisms using rank plot analysis may provide new insight into how sex-dimorphic gene expression is controlled by pulsatile somatotropic signaling in mice.

In conclusion, we propose a method that facilitates the study of GH release patterns in mice. This method provides insight into key aspects of GH release and is helpful in those small animals where obtaining serial data via cannulation is problematic. Indeed, rank plot analysis makes a valid statement about circadian endogenous GH levels, while serial data from mice reveal peak frequency. Using rank plots, we explored the acute effects of stress on circulating GH and the effects of GH on downstream signaling pathways. We also showed that extended periods of trough GH are associated with extended lifespan in mice, providing further evidence for the critical role of GH signaling in aging and longevity determination. It is noteworthy to say that other pulsatile hormones can also be studied using rank plot analysis.

**Supplementary data**

This is linked to the online version of the paper at http://dx.doi.org/10.1677/JOE-10-0317.

**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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**Author contribution statement**

MH designed, AJB, CDME, JD, MH, and LK performed the experiments. JX analyzed the data and wrote the manuscript. SR and IAM performed statistical evaluation. All authors discussed the results and commented on the manuscript.
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