

LVR-Klinikverbund



# How to Bridge the Gap between Research and Clinical Practice Examples from Anorexia Nervosa Research

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#### Conflicts of Interest

- Editor-in-chief of European Childhood and Adolescent Psychiatry
- Non-voting member of the ESCAP board
- Vice President of the European Association for the Study of Obesity
- Funding: DFG, BMBF, EU, NRW

#### Core Phenotype of Anorexia Nervosa

- Special features
  - Comparatively rather homogeneous clinical symptomatology
  - Circumscribed age manifestation range
  - Moderate to high heritability
  - Low prevalence despite ubiquitous drive for thinness
  - Historic case reports (?)
- State and trait markers
- "... the intertwining of the primary behaviors with the psychological and somatic consequences of starvation represent the core symptomatology of AN."

#### **Overview**

- Do not hesitate to ask questions: Improving the weight criterion
- Make use of spin-offs: Systematic aspects of body weight regulation in anorexia nervosa
- Identifying major questions: Starvation independent findings
- Carpe diem: Neurobiology of starvation
  - Leptin as a central switch
- Struggle to make your opinions known: Discussion of current diagnostic criteria
- Identify your role: Genetic aspects

#### DSM-IV TR Criteria for Anorexia Nervosa

- A. Refusal to maintain body weight at or above a minimally normal weight for age and height (e.g., weight loss leading to maintenance of body weight less than 85% of that expected; or failure to make expected weight gain during period of growth, leading to body weight less than 85% of that expected).
- B. Intense fear of gaining weight or becoming fat, even though underweight.
- C. Disturbance in the way in which one's body weight or shape is experienced, undue influence of body weight or shape on selfevaluation, or denial of the seriousness of the current low body weight.
- D. In postmenarcheal females, amenorrhea, i.e., the absence of at least three consecutive menstrual cycles. (A woman is considered to have amenorrhea if her periods occur only following hormone, e.g., estrogen, administration.)

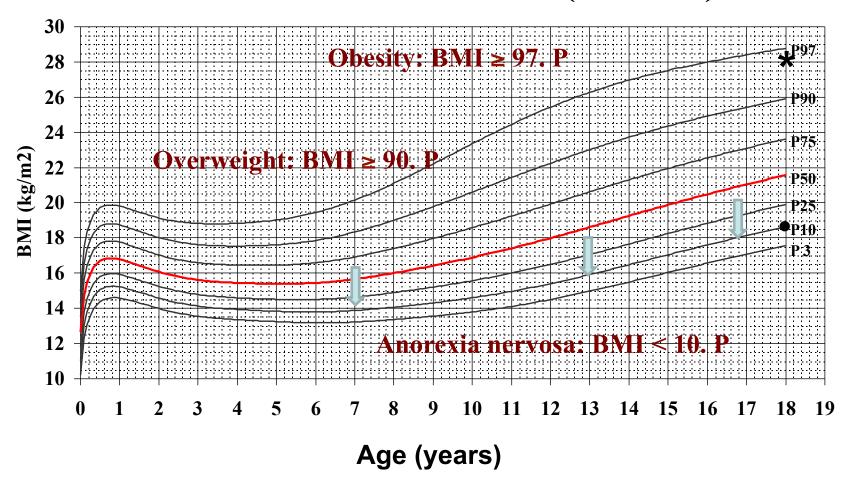
Do not hesitate to ask questions

# Absolute BMI Values Corresponding to 10<sup>th</sup> BMI Centile (NHANES I)

		Age in years						
	10-12	13-14	15-16	18-20	21-23	24-26	27-29	33-35
Fem	ales 15.6	16.6	17.4	18.4	18.5	18.5	18.7	19.4
Mal	es 15.3	16.6	17.8	19.7	20.0	20.2	20.5	21.3

Hebebrand et al., Int J Eat Dis 1996

#### **German BMI Percentiles (Females)**



Kromeyer-Hauschild et al. (2001); Monatsschrift Kinderheilkunde 149: 807

#### **US BMI Percentiles (Females)**

2 to 20 years: Girls Body mass index-for-age percentiles

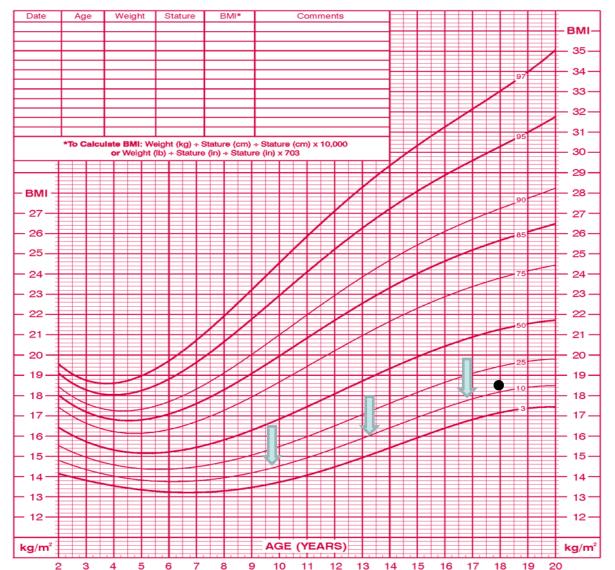
CDC

2000

growth

curves

NAME \_\_\_\_\_\_



10<sup>th</sup>
Percentile

Published May 30, 2000 (modified 10/16/00). SOURCE: Developed by the National Center for Health Statistics in collaboration with the National Center for Chronic Disease Prevention and Health Promotion (2000)



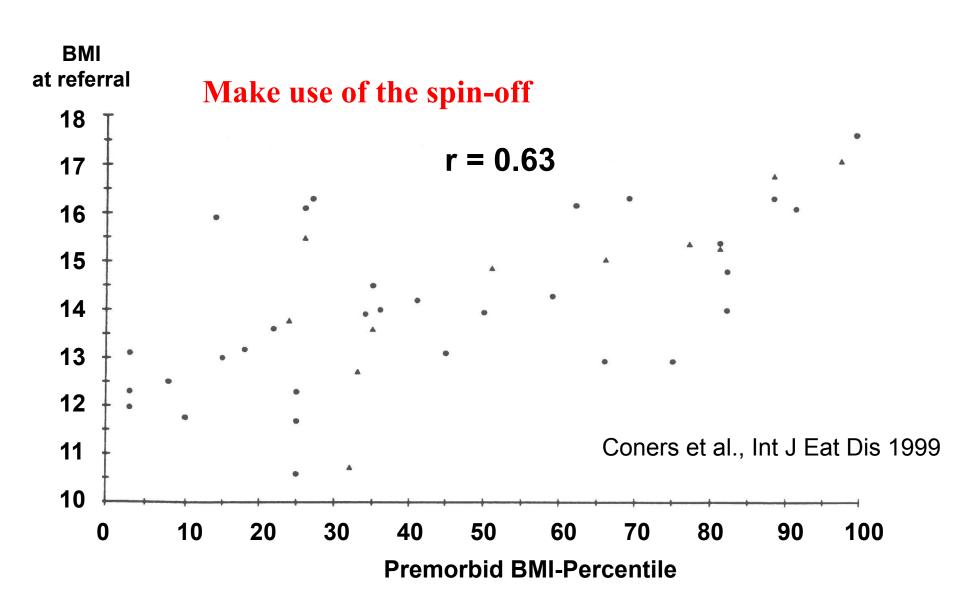
#### US Prevalence rates of BMI < 18.5 kg/m<sup>2</sup>

Table 3: Age-adjusted percent distribution (with standard errors) of underweight (BMI < 18.5 kg/m²) for adults 18 years of age and over: United States, average annual, 2005–2007 60

Age range	Males	Females
18-24 years	2.9 (0.46)	4.8 (0.40)
25–44 years	0.6 (0.08)	2.7 (0.17)
45–64 years	0.5 (0.07	1.7 (0.16)
65–74 years	1.0 (0.19)	1.9 (0.25)
75 years and over	2.4 (0.42)	4.4 (0.36)

BMI was computed using respondent-reported height and weight, without shoes.

#### Referral and Premorbid BMI



#### Referral-BMI and Weight Loss



Weight loss in kg/m<sup>2</sup> r = 0.66Coners et al., Int J Eat Dis 1999 **Premorbid BMI Percentile** 

### Relationship between Referral and Follow-up BMI: 272 Patients

#### Referral-BMI

< 13 kg/m<sup>2</sup>

≥ 13 kg/m<sup>2</sup>

n = 100

n = 172

#### BMI at follow-up (mean: 9.5 years)

 $\leq$  17.5 kg/m<sup>2\*</sup>

35%

**12.8%** 

≤ 5<sup>th</sup> centile\*

44%

19.8%

≤ 10<sup>th</sup> centile\*

**56%** 

29.0%

 $\geq$  25 kg/m<sup>2</sup>

1%

3.4%

**Deceased** 

11%

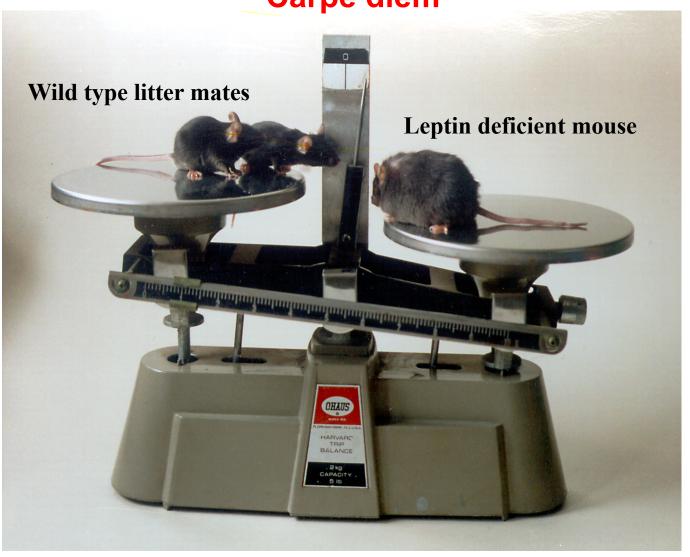
0.6%

Hebebrand et al., Am J Psychiatry 154: 566-569; 1997

<sup>\*</sup>including deceased patients

#### The *ob/ob* mouse

Carpe diem



Zhang et al., Nature 1994

#### courtesy of Sadaf Farooqi



Child B before leptin

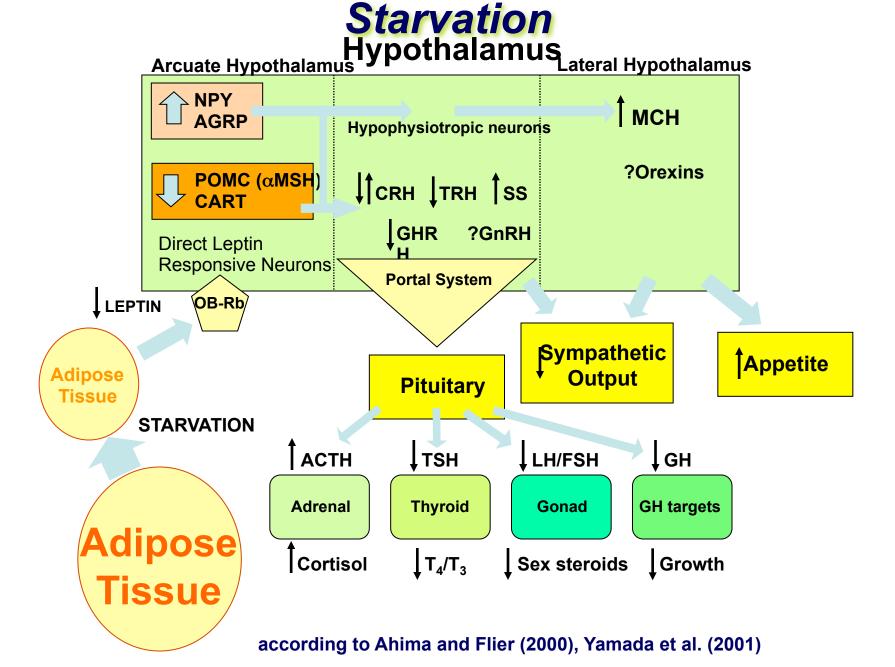
 $(wt = 42kg \ at \ 3yrs)$ 



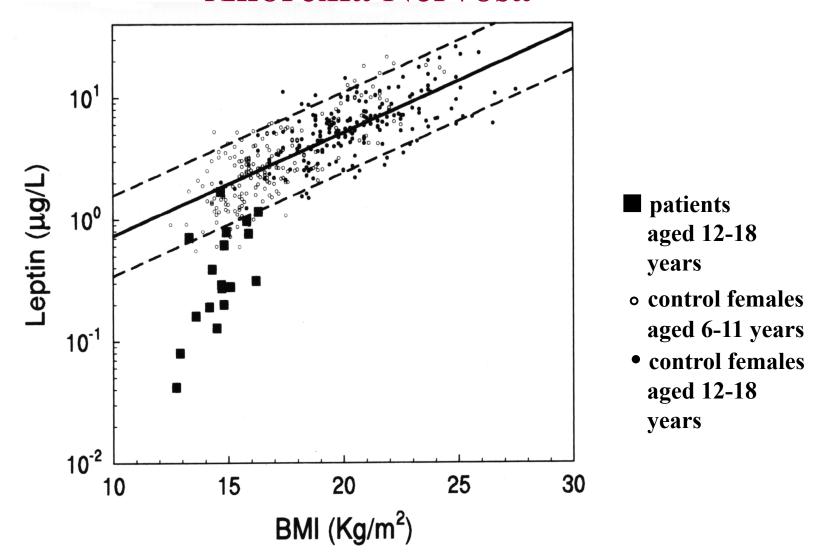
Child B after leptin

 $(wt = 32kg \ at \ 7yrs)$ 

#### Leptin Signaling: Adaptation to Semi-



#### Serum lg10 Leptin Levels in Acute Anorexia Nervosa



Hebebrand et al 1995, 1997; The Lancet, Mol Psychiatry

#### Evidence for a leptin threshold in AN

#### Central hypothesis:

Hypoleptinemia in AN patients; healthy underweight females have higher levels

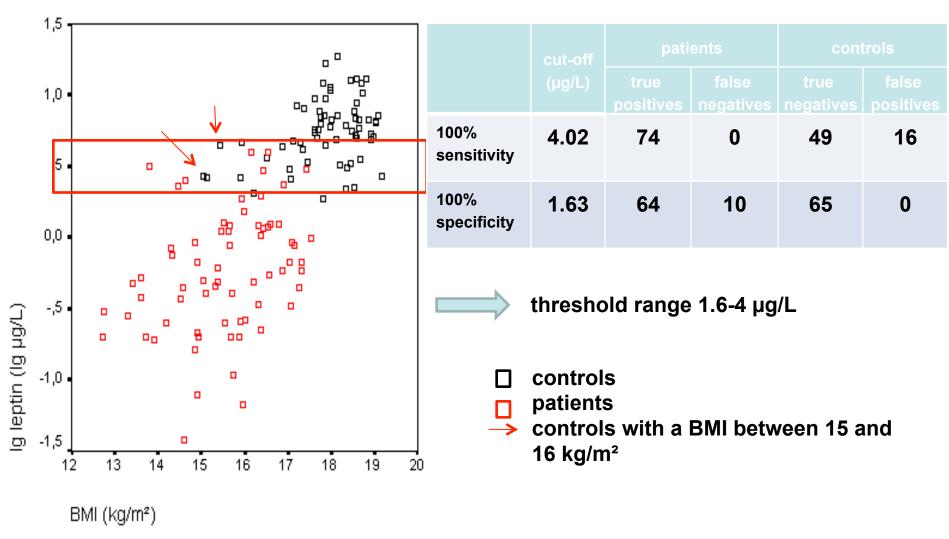


existence of a **threshold** value/range

#### Background:

- hypoleptinemia: a cardinal feature of semistarvation in AN
- according to most studies leptin levels only infrequently exceed
   μg/L in patients with AN (Müller et al. 2009)
- threshold of 1.85µg/L separates patients with AN from healthy underweight females (Köpp et al., 1997)

### Hypoleptinemia: Sensitivity and Specificity for Diagnosis of Anorexia nervosa



Föcker et al. 2011, J Neural Transmission

## Hypoleptinemia: A biological marker for Anorexia nervosa

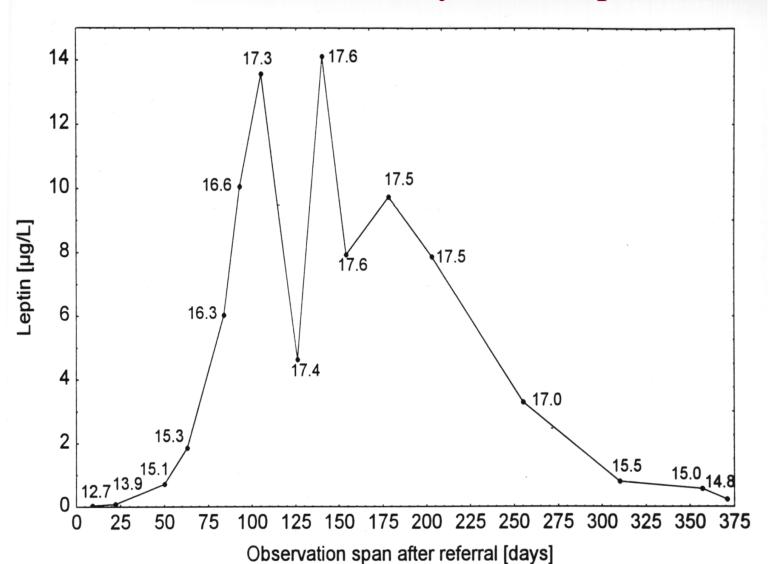
cut off (μg/L)	sensitivity	specificity	ppV	npV
2.0	0.89	0.97	0.604	0.994
2.5	0.93	0.94	0.444	0.996

statistical paramter	legend
positive predictive Value	% of individuals with positive test results who are correctly diagnosed as patients
negative predictive Value	% of individuals with negative test results who are correctly diagnosed as healthy
sensitivity	% of actual positives which are correctly identified as such
specificity	% of negatives which are correctly identified



a threshold in the range of 2 µg/L appears appropriate for screening purposes

Serum leptin levels in a patient with anorexia nervosa over a one year time period



Hebebrand et al 1998; Mol Psychiatry

#### Leptin and Anorexia Nervosa

 Assessment of the clinical implications of hypoleptinemia and hyperleptinemia

 Anorexia nervosa is a model disorder to assess the effects of hypoleptinemia and of the rapid transition from hypoleptinemia to hyperleptinemia in humans

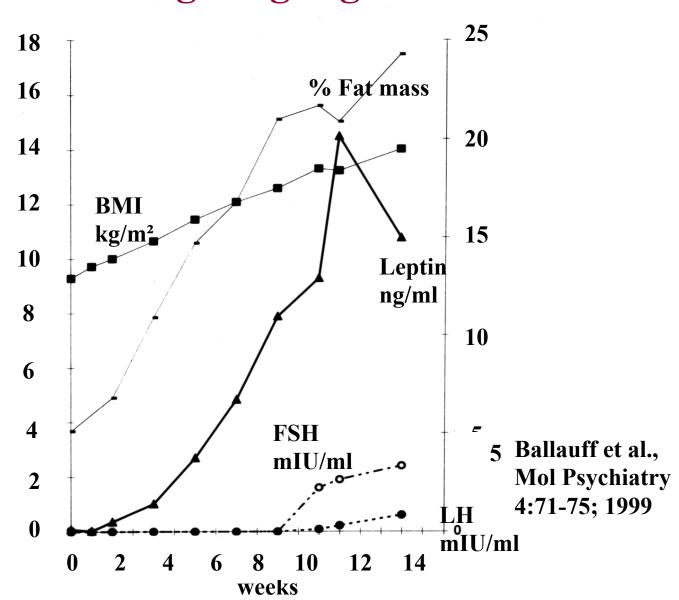
# Semi-starvation in humans: selected physical and laboratory findings

amenorrhea hypothermia cold intolerance hypotension bradycardia dryness of skin lanugo constipation abdominal pain increased ventricular -brain ratio

reduced FSH, LH, estrogen low T3 syndrome high ghrelin levels reduced hematopoiesis hyperadrenocorticism reduced resting energy hypoleptinemia

APA, 1994 Hebebrand et al., 1995, 1997 Otto et al., 2001

# Serum leptin and gonadotropin levels during weight gain



# Hypothalamic Amenorrhea: Treatment with Leptin

- 8 females with hypothalamic amenorrhea of ≥ 6 months duration (mean: 5 years)
- 6 untreated controls
- Leptin treatment (r-metHuLeptin) for three months
  - Increment of serum LH levels within 2 weeks
  - Increase of the maximal follicle diameter, size of the ovary and increment of serum estrogen level within 3 months
  - 3 patients ovulated, 2 pre-ovulatory follicles
  - No significant weight loss; no side effects except reduction of appetite in third month of treatment

Welt et al., N Engl J Med 351: 987-97, 2004

#### Influence of leptin on brain growth

- Leptin treatment of *ob/ob* mice increases weight of brain
- 3 adults with leptin deficiency treated with recombinant leptin
- MRI at baseline, 3, and 18 months after initiation of treatment
- Volume increments of gray matter in frontal gyrus cinguli, inferior parietal lobe and cerebellum

# Semi-starvation in humans: psychological findings

depressed mood
social withdrawal
pre-occupation with food
rigidity
hunger
abnormal eating behavior
reduced libido

irritability
inflexible thinking
limited spontaneity
restrained initiative
restrained emotional
expression
loss of ambition

Keys et al., 1950; APA, 1994

#### **Hyperactivity and Anorexia Nervosa:**

different aspects of the phenomenon and clinical terminology

#### Identify good questions

- Hyperactivity / elevated activity
- Motor restlessness / diffuse or nocturnal restlessness
- Excessive or extensive physical activity / intensive engagement in sports
- Compulsive physical activity, exaggerated need of physical activity
- Behavioral activation, paradoxical liveliness, excessive vitality, surplus of physical energy

Prevalence rates: 30% - 80%

#### Factors Associated with "Hyperactivity" in Anorexia Nervosa

- Early onset
- High physical activity levels during childhood
- Inverse correlation between food intake and level of physical activity during the acute stage of the disorder
- Reduction of the inner restlessness during therapeutically induced weight gain
- Correlation with anxiety, irritability and obsessive-compulsive symptoms
- Worse prognosis

# Semi-starvation induced hyperactivity

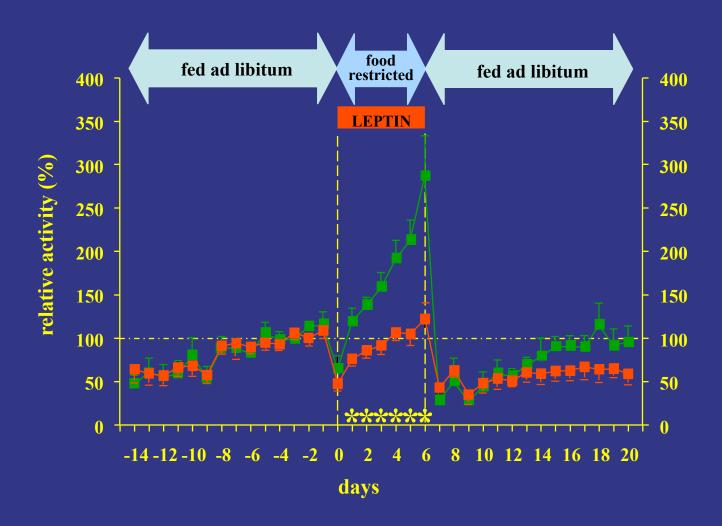
#### **Anorexia based hyperactivity**

In rats caloric restriction leads to semi-starvation induced hyperactivity.

Model for anorexia nervosa?

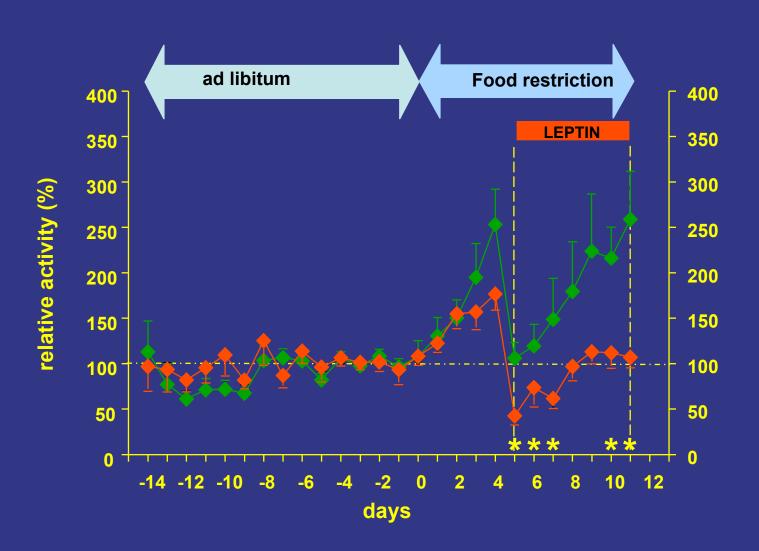


#### **Leptin Suppresses Semi-Starvation Induced Hyperactivity**

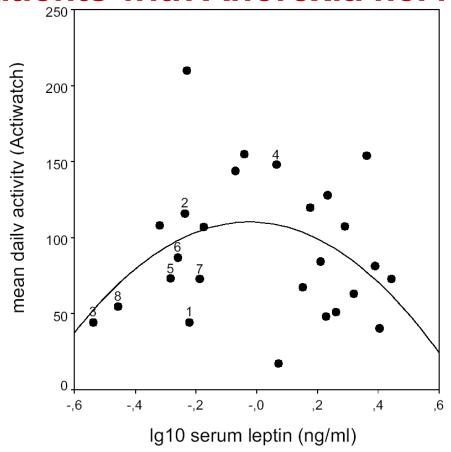


Exner et al., Mol Psychiatry 5: 476-481, 2000

#### "Treatment" of Hyperactivity



## Serum Leptin Levels and Mean Daily Activity in Patients with Anorexia nervosa



**Figure 1:** Scatterplot of mean daily physical activity of 72 hours (Actiwatch output) vs. lg10 serum leptin levels of 26 patients with anorexia nervosa. Partial correlation controlled for BMI: r= -.410, p=.042. Numbers 1-8 indicate the patients with lowest BMI within the study sample.

Holtkamp et al., 2006; Biol Psychiatry 60:311-3

# Activity in Semi-Starvation: Minnesota Study

• The attitude of the men to physical exertion was ambivalent. It made them tired and as a rule was avoided. On the other hand, occasionally some men exercised deliberately. Thus certain subjects attempted to lose weight by driving themselves through periods of excessive expenditure of energy with the object of either obtaining increased bread rations (when weight loss exceeded the prescribed rate) or avoiding reduction in rations (when weight loss lagged)

subjects moved slowly and cautiously

curtailment of spontaneous activity

coordination was affected

Franklin, Schiele, Brozek and Keys, J Clin Psychol 4: 28-45, 1948

• the men rated themselves as ..... restless, unable to concentrate, and markedly "nervous"

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- D. In postmenarcheal females, amenorrhea, i.e., the absence of at least three consecutive menstrual cycles. (A woman is considered to have amenorrhea if her periods occur only following hormone, e.g., estrogen, administration.)

Struggle to make your opinions known or don't give up

# Refusal to Maintain Body Weight at or above a Minimally Normal Weight?

- Refusal evidently implies an active, conscious <u>and</u> willful psychological process
- But: Evidence for underlying regulatory phenomena contributing to both somatic and mental symptoms and the course of disorder
  - Weight course a non-random process
  - Hypoleptinemia underlies amenorrhea
  - Hypoleptinemia contributes to hyperactivity
  - Hyperleptinemia predicts relapse
  - Genetics

# Refusal to Maintain Body Weight at or above a Minimally Normal Weight?

- Patients do seek help
- Inferred behavior instead of description of behavior
- Term refusal is not used for any other psychiatric disorder
  - Anxiety, affective, conduct disorders
- Term refusal can be perceived as conveying a paternalistic and prejorative attitude
- No systematic evidence for the term
- Refusal not assessed in standard diagnostic interviews

#### DSM-5 criteria for Anorexia nervosa

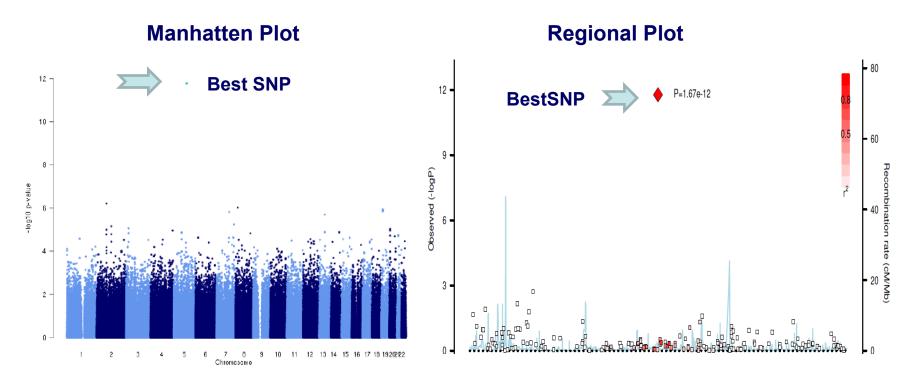
- 1. Restriction of energy intake relative to requirements leading to a significantly low body weight in the context of age, sex, developmental trajectory, and physical health.
- 2. Intense fear of gaining weight or becoming fat, even though underweight.
- 3. Disturbance in the way in which one's body weight or shape is experienced, undue influence of body weight or shape on self-evaluation, or denial of the seriousness of the current low body weight.

#### **GWAS** for Anorexia Nervosa



### International Multicenter Study funded by ,Welcome Trust Case Control Consortium (WTCCC3)

- Coordinators: C. Bulik (Chapel Hill, USA) and D. Collier (London/UK)
- n= 2,907 AN patients (n= 500 from Germany) and 14,860 controls (Illumina 660W-Quad)



#### **Identify your role**

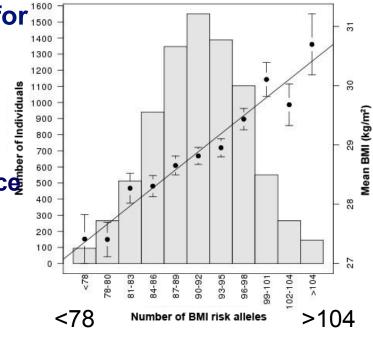
#### **GIANT: BMI**



Genetic Investigation of **ANthropometric Traits** 

Meta-analysis (GWAS and Metabochip) for 1500 **BMI** 

- ≤ 339,224 individuals
- 97 BMI loci (56 novel)
- 2.7% of BMI variance explained
- BMI loci (56 novel)
  % of BMI variance explained
  Frequent alleles explain ≤ 20% of variance
- **Role of CNS**



Relevant pathways: e.g. synaptic function, glutamate signaling, insulin secretion/ action, energy metabolism, lipid biology

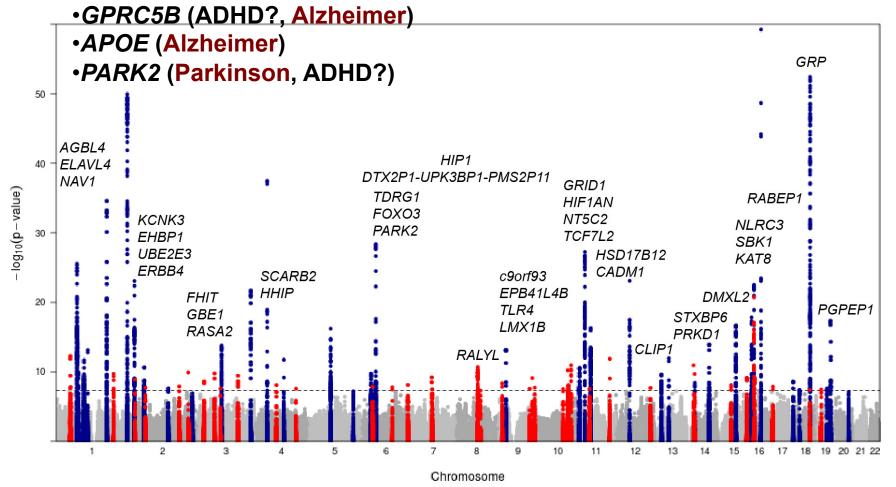
#### **GIANT: BMI**



Genetic Investigation of ANthropometric Traits

#### Genes with potential relevance for neuropsychiatric disorders:

•BDNF (ADHD?, MDD?)



Locke et al. 2015, Nature. 2015 Feb 12;518:197-206

# Three Loci Potentially Involved in both Anorexia Nervosa and Obesity

Look-up of the 1000 SNPs with lowest p-values of a GWAS for AN (Boraska et al, 2014) in GWAS meta-analysis for BMI variation (Locke et al, 2015)

Significant association (p-values < 5x10<sup>-05</sup>, Bonferroni corrected p < 0.05) for 9 SNPs at 3 independent loci (chr. 2, 10 and 19)

All risk alleles were directionally consistent for AN and obesity

# Leptin hebt inhibitorischen Effekt einer Futterrestriktion auf das Längenwachstum auf

- Das Längenwachstum von jungen futterrestriktierten Mäusen ist beeinträchtigt
- Leptin unterdrückt den wachstumshemmenden Effekt einer Futterrestriktion (Tibia)
  - Leptin wirkt über Leptinrezeptoren in Wachstumsfugen

#### Acknowledgements

Clinical work: Anne Ballauf, Anke Hinney, Helmut Remschmidt, Christian Holtkamp, Beate Herpertz-Dahlmann

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Criteria: Cindy Bulik, Regina Casper, Ulrich Schweiger, Janet Treasure