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SCIENTIFIC DATA

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SUBJECT CATEGORIES

- » Non-alcoholic fatty liver disease
- » Metabolomics
- » Gene expression analysis
- » Systems biology

Multi-omic profiles of human non-alcoholic fatty liver disease tissue highlight heterogenic phenotypes

Wasco Wruck¹, Karl Kashofer², Samrina Rehman³, Andriani Daskalaki⁴, Daniela Berg⁵, Ewa Gralka⁶, Justyna Jozefczuk⁴, Katharina Drews⁴, Vikash Pandey⁴, Christian Regenbrecht⁷, Christoph Wierling⁴, Paola Turano⁶, Ulrike Korf⁵, Kurt Zatloukal², Hans Lehrach⁴, Hans V. Westerhoff^{3,8,9} & James Adjaye^{1,4}

Non-alcoholic fatty liver disease (NAFLD) is a consequence of sedentary life style and high fat diets with an estimated prevalence of about 30% in western countries. It is associated with insulin resistance, obesity, glucose intolerance and drug toxicity. Additionally, polymorphisms within, e.g., *APOC3*, *PNPLA3*, *NCAN*, *TM6SF2* and *PPP1R3B*, correlate with NAFLD. Several studies have already investigated later stages of the disease. This study explores the early steatosis stage of NAFLD with the aim of identifying molecular mechanisms underlying the etiology of NAFLD. We analyzed liver biopsies and serum samples from patients with high- and low-grade steatosis (also pre-disease states) employing transcriptomics, ELISA-based serum protein analyses and metabolomics. Here, we provide a detailed description of the various related datasets produced in the course of this study. These datasets may help other researchers find new clues for the etiology of NAFLD and the mechanisms underlying its progression to more severe disease states.

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Design Type(s)	parallel group design • transcription profiling by array design • protein expression profiling
Measurement Type(s)	transcription profiling assay • protein expression profiling • metabolite profiling
Technology Type(s)	microarray platform • ELISA • nuclear magnetic resonance assay
Factor Type(s)	disease stage • obesity
Sample Characteristic(s)	Homo sapiens • serum • liver

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Background & Summary

With an estimated prevalence of about 30% in western countries, NAFLD is a major public health issue¹. Sedentary life-style and excessive food consumption correlate with rate at which NAFLD cases appear. Epidemiologic studies showing a prevalence of the disease that differs between countries as well as between groups in the same country, appear to reflect an interplay of environmental and genetic factors in its etiology¹. Additionally, polymorphisms in, e.g., *APOC3*, *PNPLA3*, *NCAN*, *TM6SF2* and *PPP1R3B*, correlate with NAFLD^{2,3}. Over-feeding directly induces insulin resistance⁴. Causality between steatosis and the metabolic syndrome of insulin resistance, obesity, and glucose intolerance, is still unresolved⁵. While the correlation between steatosis and insulin resistance is established there is debate about the relationship between steatosis and hepatic insulin resistance⁶. Samuel *et al.* showed that activated *PKC-ε* and *JNK* can induce insulin resistance via impaired *IRS1* and *IRS2* tyrosine phosphorylation in rats fed with high fat diet⁷. An investigation on the insulin-like growth factor (IGF) axis in the Nurses' Health Study⁸ and another population study of 3863 people⁹ addressed connections between the IGF axis, insulin resistance, diabetes risk and NAFLD. *IGFBP3* is associated with various cancers and up-regulation of *IGF1* receptor (*IGF1R*) is considered an early event in hepatocarcinogenesis¹⁰. Thus, the IGF axis might play an important role in a direct development of carcinoma from steatosis without the formerly assumed intermediary phase of cirrhosis¹¹.

The progression of NAFLD from mild steatosis up to severe steatohepatitis and even liver cirrhosis and hepatocellular carcinoma, varies widely between individual patients. Insulin resistance, dysregulation of cytokines as a basis for inflammation, and oxidative stress appear to foster progression to steatohepatitis¹². A two-step progression from simple steatosis to steatohepatitis and fibrosis has been proposed¹³, and suggests that after fat accumulation in the liver due to insulin resistance, lipids are peroxidized with cytokines and Fas ligand induced by excessive ROS. However, this two-step progression has been questioned⁵. We found that in fibroblasts derived from steatosis patients *AKT/mTOR* signaling was reduced and that the insulin-resistant phenotype is exhibited not only by insulin-metabolizing central organs, e.g., the liver, but also by skin fibroblasts¹⁴. Transcriptome data identified a regulatory network orchestrated by the transcription factor *SREBF1* and linked to a metabolic network of glycerolipid and fatty acid biosynthesis. The downstream transcriptional targets of *SREBF1* which include the phosphatidic acid phosphatase *LPIN1* and *LDLR*, were also involved.

Moreover, there is the possible involvement of ROS in disease progression. Houstis *et al.*¹⁵ demonstrated that oxidative stress can induce insulin resistance and that anti-oxidants may ameliorate insulin resistance. Depletion of glutathione can improve insulin sensitivity in mice¹⁶. Glutathione is known as the body's master antioxidant, protecting cells against damage caused by numerous reactive intermediates¹⁷. Detoxification of these reactive metabolites results in the consumption of glutathione either via oxidation or conjugation. Maintenance of the intracellular glutathione level is thereby a critical liver function, which could be impaired following insult/injury or in steatosis and steatohepatitis.

Several other studies exploring various aspects of NAFLD have been published. A recent publication by Moylan *et al.* showed that it is possible to discriminate mild versus severe fibrosis stages of NAFLD patients via their gene expression profiles¹⁸. Another study from Speliotes *et al.* investigated NAFLD via a genome-wide association study (GWAS) approach³. Besides the most prominent association of *PNPLA3* this study reported several other associations including one at locus 19p13.11 which is in strong linkage disequilibrium with a recently found steatosis-linked polymorphism in *TM6SF2*, transmembrane6 superfamily member 2 (refs 19,20). A knockdown of *TM6SF2* in human hepatoma cell lines and in mice led to an increase in lipid droplet area while overexpression led to a decrease¹⁹.

Interestingly, the above mentioned genes associated with NAFLD in GWAS were not detected in a large-scale GWAS about obesity and insulin biology although the metabolic syndrome connects NAFLD and obesity²¹. Feldstein *et al.* found CK-18 as a non-invasive biomarker for NASH by comparison of plasma samples from patients with biopsy proven NAFLD²². Du Plessis *et al.* used analysis results from subcutaneous and visceral fat and liver biopsies to construct a model which predicts NAFLD liver histology²³. This model involves the genes *CCL2*, *DMRT2*, *GADD45B*, *IL1RN*, and *IL8*. In contrast to the studies of Moylan *et al.* and Feldstein *et al.* our study highlights potential means of classifying distinct grades of Steatosis in NAFLD—the very early stage of the disease. Although it is evident that a complex interplay of genetic and environmental factors contribute to the development of steatosis, to date there has not been a systemic study of the disease employing a multi-omic approach—transcriptome, ELISA-based proteome and metabolome. Therefore, the intention of this study is to provide a more comprehensive view of steatosis based on transcriptomic, metabolomic and protein biomarker profiles. Additionally, this should lay down the foundation for follow-up systems biology-based studies.

In the current study we analyzed patient liver biopsies and associated serum samples, from patients with the insulin resistance phenotype confirmed by the HOMA-IR model²⁴. Here, we describe these valuable data sets deposited in public repositories, which might support other researchers in identifying new clues for the etiology of NAFLD and the mechanisms underlying its progression to more severe disease states.

ID	gender	Age	BMI	% steatosis	grouping by pathologist	steatosis grouping	medical centre	liver illumina array rep.1 (GSE46300)	liver illumina array rep.2 (GSE46300)	serum NMR data	serum ELISA data
H0004	f	54	47	10%		obese, low steatosis	Graz (Austria)	GSM1128362	GSM1128363	MTBLS174	10.6084/m9.figshare.1333564
H0007	f	33	51	40%		obese, high steatosis	Graz (Austria)	GSM1128364	GSM1128365	MTBLS174	10.6084/m9.figshare.1333564
H0008	m	61	46	40%	obese, high steatosis	obese, high steatosis	Graz (Austria)	GSM1128366	GSM1128367	MTBLS174	10.6084/m9.figshare.1333564
H0009	f	48	49	5–10%	obese, low steatosis	obese, low steatosis	Graz (Austria)	GSM1128368	GSM1128369	MTBLS174	10.6084/m9.figshare.1333564
H0011	f	58	45	70%	obese, high steatosis	obese, high steatosis	Graz (Austria)	GSM1128370	GSM1128371	MTBLS174	10.6084/m9.figshare.1333564
H0012	f	50	35	0	obese, low steatosis	obese, low steatosis	Graz (Austria)	GSM1128372	GSM1128373	no	no
H0018	f	35	41	30–40%	obese, high steatosis	obese, high steatosis	Graz (Austria)	GSM1128374	GSM1128375	MTBLS174	10.6084/m9.figshare.1333564
H0021	m	49	41	0%		no steatosis	Graz (Austria)	GSM1128376	GSM1128377	MTBLS174	10.6084/m9.figshare.1333564
H0022	m	45	49	40%		obese, high steatosis	Graz (Austria)	GSM1128378	GSM1128379	MTBLS174	10.6084/m9.figshare.1333564
H0024	m	29	44	50%		obese, high steatosis	Graz (Austria)	no	no	MTBLS174	10.6084/m9.figshare.1333564
H0025	f	53	46	15–20%		obese, low steatosis	St Gallen (Switzerland)	no	no	MTBLS174	10.6084/m9.figshare.1333564
H0026	f	46	39	0%		no steatosis	St Gallen (Switzerland)	no	no	MTBLS174	10.6084/m9.figshare.1333564
H0027	m	44	42	50%		obese, high steatosis	St Gallen (Switzerland)	no	no	MTBLS174	10.6084/m9.figshare.1333564
H0028	f	28	43	20%		obese, low steatosis	St Gallen (Switzerland)	no	no	MTBLS174	10.6084/m9.figshare.1333564
H0029	f	40	39	< 5%		no steatosis	St Gallen (Switzerland)	no	no	MTBLS174	10.6084/m9.figshare.1333564
H0030	m	22	45	30%		obese, low steatosis	St Gallen (Switzerland)	no	no	MTBLS174	10.6084/m9.figshare.1333564
H0031	m	22	41	0%		no steatosis	St Gallen (Switzerland)	no	no	MTBLS174	10.6084/m9.figshare.1333564
H0033	f	44	43	40%		obese, high steatosis	St Gallen (Switzerland)	no	no	MTBLS174	10.6084/m9.figshare.1333564
H0034	m	50	42	10%		obese, low steatosis	St Gallen (Switzerland)	no	no	MTBLS174	10.6084/m9.figshare.1333564

Table 1. Samples related to data sets in repositories (Data Citations 1–Data Citation 3).

Methods

Patient recruitment, sample collection and clinical measurements

All patients participating in this study were recruited in the Multidisciplinary Obesity Research (MORE) project at the Medical University of Graz, Austria or at the Interdisciplinary Adipositas Center at the Kantonsspital St Gallen, Switzerland. Patients with morbid obesity who admitted into hospital for treatment by bariatric surgery (gastric banding, gastric bypass, sleeve gastrectomy) were invited to participate in the study and to sign the informed consent. The study was approved by the institutional review board of the Medical University of Graz (reg. IRB00002556 at the Office for Human Research Protections of the US Departments of Health and Human Services) under license 20–143 ex 08/09. All experiments were performed in accordance with approved guidelines. Written informed consent was obtained from all participants. In the course of the bariatric surgery, samples of blood, skin and a liver biopsy were taken. Out of 18 patients (Table 1), 9 liver biopsies were of high quality enabling their use in the transcriptome analyses. Serum plasma was available from all the patients. The overall experimental design of this study is illustrated in Fig. 1. A pathological diagnosis of the liver phenotype, including liver steatosis grading based on H&E morphology, was performed by an experienced, board certified pathologist (CL). We simplified Kleiner's scoring scheme by condensing Steatosis grades 2 (34–66%) and 3 (> 66%) to our 'high-grade' while adopting grades 0 ('none') and 1 ('low')²⁵. This simplification was made because the inter-patient-variability in this complex heterogeneous disease did not allow a more detailed grading on the omics levels. Two examples of liver biopsies are shown in Fig. 2a.

Illumina bead chip hybridization and data analysis

Microarray experiments were carried out on the Illumina BeadStation 500 platform (Illumina, San Diego, CA, USA). Briefly, 500 ng DNase-treated total RNA were used as input for amplification and biotin labeling reactions (Illumina TotalPrep RNA Amplification Kit, Ambion) prior to hybridization of the resulting cRNAs onto Illumina HumanHT-12_v4_BeadChips, washing, Cy3-streptavidin staining and scanning according to the manufacturer's instructions.

Transcriptomics data analysis

Illumina data was processed via R/Bioconductor²⁶ and packages lumi²⁷, limma²⁸ and biomaRt. Background-corrected log₂-transformed data was normalized via quantile normalization from the lumi package.

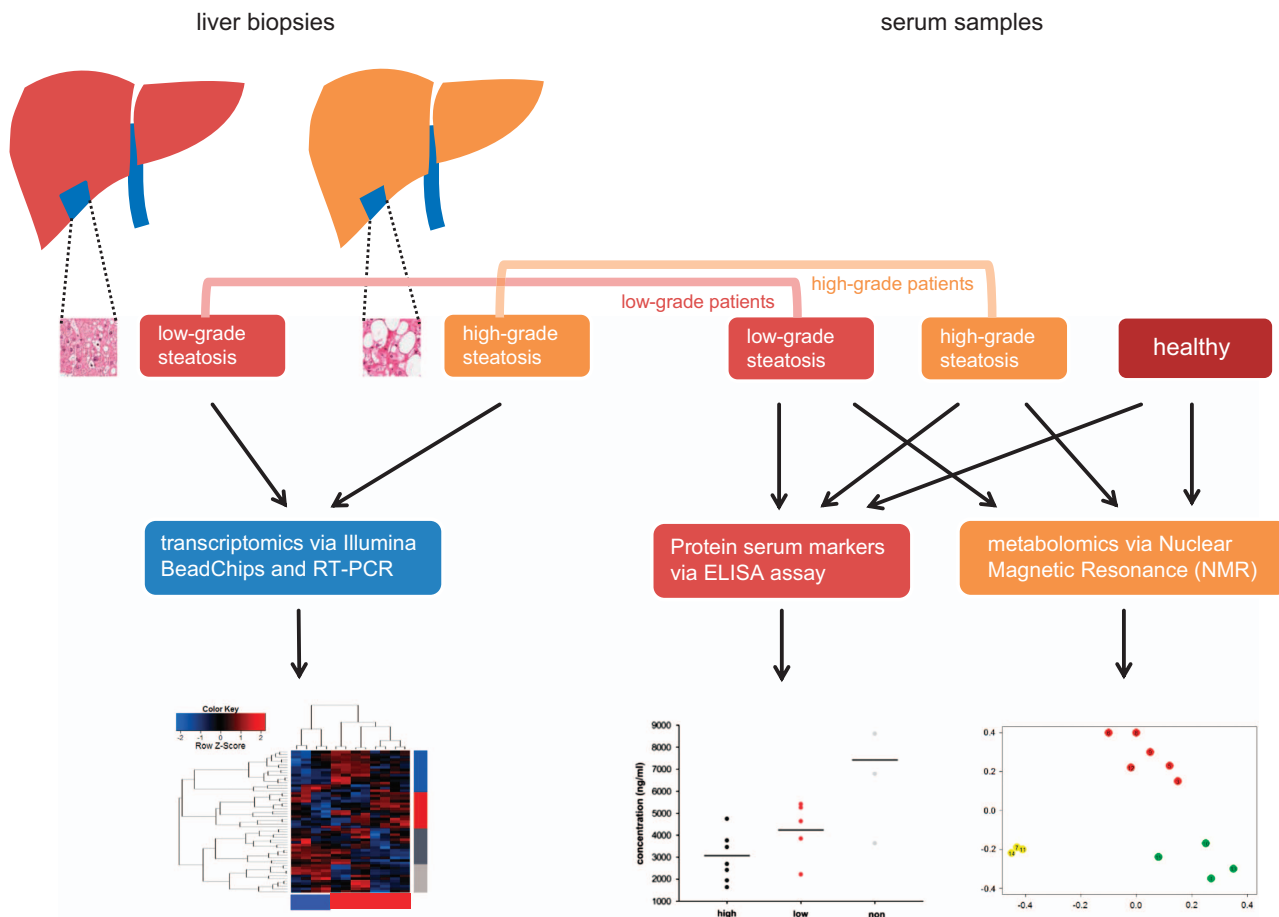


Figure 1. Scheme of experiments for multi-omics comparison of steatosis grades. The scheme shows how the distinct severities of non-alcoholic fatty liver disease (NAFLD) are compared in terms of transcriptomics, metabolomics and potentially relevant parts of the proteome. Liver biopsies were taken from NAFLD patients and classified by pathologists as low-grade (5–33% steatosis area) and high-grade (>33% steatosis area). The transcriptome of liver biopsies were assessed on Illumina HumanHT-12 v4 BeadChips and on RT-PCR. Serum samples of these NAFLD patients and from healthy persons were taken and investigated at the protein level employing ELISA assays and at the metabolome level via Nuclear Magnetic Resonance (NMR).

qRT-PCR

Quantitative real-time polymerase chain reaction (qRT-PCR) was carried out to confirm the microarray-derived data. Reactions were carried out on the ABI PRISM 7900HT Sequence Detection System (Applied Biosystems). Data analysis was carried out using the ABI PRISM SDS 2.2.1 software (Applied Biosystems) and Microsoft Excel (Microsoft Corporation, Redmond, WA, USA). GAPDH-normalized, relative mRNA levels of each gene (high steatosis versus low steatosis) were calculated based on the $2^{-\Delta\Delta CT}$ Method. Primer sequences for QRT-PCR validation are described in Table 2.

ELISA-based assay for biomarkers

ELISA measurements from plasma samples were carried out using the Ciraplex platform (Aushon Biosystems, Billerica, MA, US). Commercial assays were purchased and measurements were carried out according to instructions provided by the manufacturer. The following 29 targets were analyzed either as single-plex assays or as multiplex assay: hFGFb; hGROa; hLIF;hIFN γ ; hIL1b; IL4; IL5; hIL6; hIL10; hIL12p70; hIL13; hTNFa; hI309; hIL8; hIP10; hMCP4; hMIP1a; hMIP1b; hCRP; hLeptin; hPAIactive; hResistin; hIGFBP1; hIGFBP3; hIGFBP2; hMIF; hApoA1; hCRP; hAcrp30.

NMR sample preparation

Frozen plasma samples were thawed at room temperature and shaken before use. According to standard methodologies a total of 300 μ l of buffer (70 mM Na $_2$ HPO $_4$; 20% (v/v) D $_2$ O; 6.15 mM NaN $_3$; 6.64 mM TMSP; pH 7.4) was added to 300 μ l of each serum sample. A total of 450 μ l of this mixture was transferred into a 4.25 mm NMR tubes (Bruker BioSpin) for analysis.

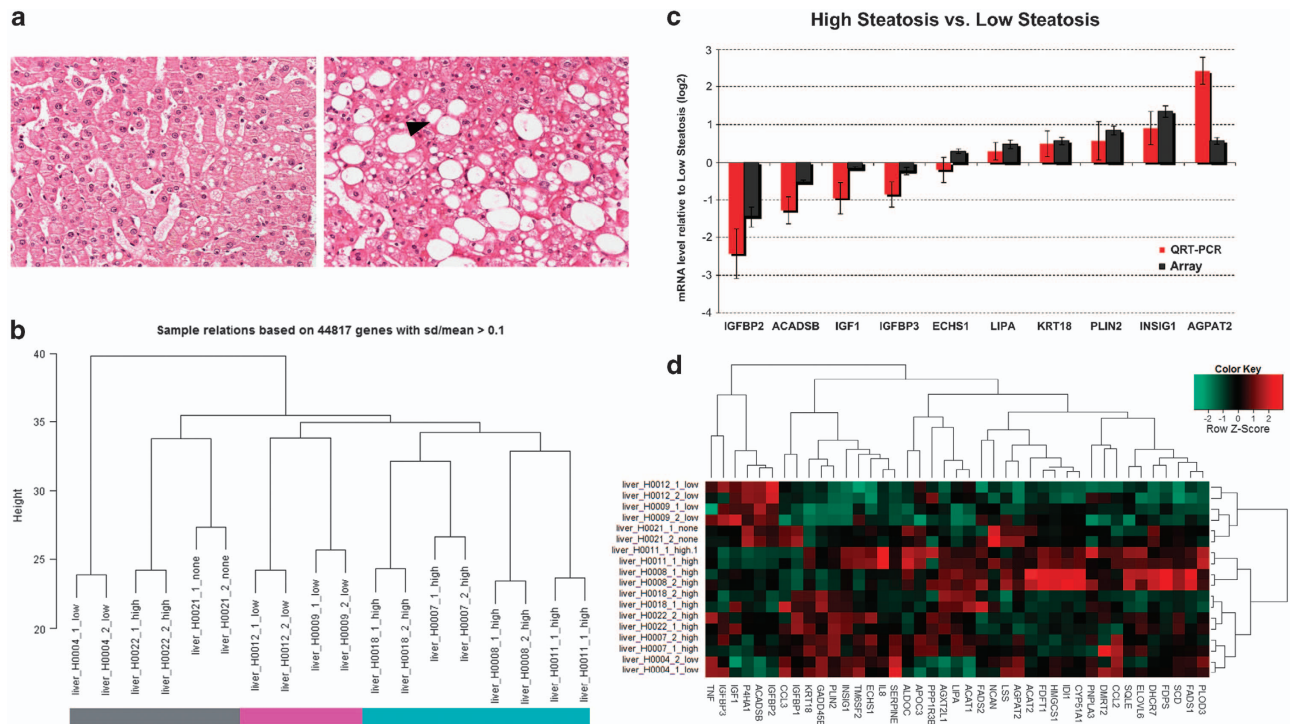


Figure 2. Histopathological and transcriptome characterization of liver tissue. **(a)** Liver tissue with only marginal pathological changes (H9, low-grade steatosis group). The hepatocytes are arranged in one cell thick plates, separated by sinusoids. They contain only few small isolated fat valvules (H&E stained section). Hepatocytes of the intermediate and central lobular areas contain macrovesicular fat (image to the right, H8, steatosis group, hepatocytes with fatty change are indicated by arrow heads; H&E stained section). **(b)** Hierarchical clustering of the transcriptomes of patient liver samples. We identified three clusters: high (>33%) steatosis (cyan), low (5–33%) steatosis (magenta) and heterogeneous clusters of high, low and no steatosis (grey). **(c)** Quantitative QRT-PCR confirmation of genes differentially expressed in high versus low steatotic livers. The columns represent the mean of four biological replicates (high steatosis) versus two biological replicates (low steatosis). Error bars indicate standard errors of the mean. Array-derived and RT-PCR-derived columns are depicted in dark grey and red respectively. **(d)** Heatmap of genes differentially expressed in high versus low steatotic livers and genes found in literature and in genome-wide association studies.

NMR spectra acquisition and processing

NMR spectra from 18 plasma samples from morbidly obese patients that underwent different type of bariatric surgery and additionally have developed steatosis were collected (Table 1). ¹H-NMR spectra were acquired using a Bruker spectrometer (Bruker Biospin). Unsupervised and supervised methods were used in order to identify a disease-related metabolomic profile that might contain a signature of steatosis.

Data Records

Data record 1

The microarray experiments discussed in this publication were carried out on the Illumina BeadStation 500 platform (Illumina, San Diego, CA, USA). The data have been deposited in NCBI's GEO and are accessible through GEO Series accession number GSE46300 (Data Citation 1).

Data record 2

Metabolomic raw data from Nuclear magnetic resonance (NMR) measurements have been deposited at the MetaboLights database (<http://www.ebi.ac.uk/metabolights>) of the European Bioinformatics Institute (EBI) under MTBLS174 (Data Citation 2).

Gene	Fwd	Rev	Product size
ACADSB	CACCATTGCAAGCATATCG	GCAAGGCACTTACTCCCAAC	117
AGPAT2	GGGGCGTCTTCTTCATCA	TTGAGGTTCTCCCTGACCAT	91
ECHS1	AACCTTTGCCACTGATGACC	CAAGCAGAGGTGTGAAGCAG	112
IGF1	TGCAGGAGGACTCTGAAAC	AGCTGCGTGATATTTGAAAGG	111
IGFBP2	CTCCCTGCCAACAGGAACCTG	TCTTGCACTGTTTGAGGTTGTACAG	147
IGFBP3	CAACTGTGGCCATGACTGAG	CCTGACTTTGCCAGACCTTC	92
INSIG1	CAACACCTGGCATCATCG	CTCGGGAAGAGAGTGACAT	118
KRT18	GAGGTGGAGCTGCTGAGAC	CAAGCTGGCCTTCAGATTTC	99
LIPA	CATCTGTGTAAGCCAAAGC	AATCCCTGAGCTGAGTTTGC	112
PLIN2	GCTGAGCACATTGAGTCACG	TGGTACACCTGGATGTTGG	102

Table 2. Primer sequences for QRT-PCR validation of genes differentially expressed between high-grade and low-grade steatosis.

sample	liver_H0007_1_high	liver_H0007_2_high	liver_H0008_1_high	liver_H0008_2_high	liver_H0011_1_high	liver_H0011_2_high	liver_H0018_1_high	liver_H0018_2_high	liver_H0022_1_high	liver_H0022_2_high	liver_H0004_1_low	liver_H0004_2_low	liver_H0009_1_low	liver_H0009_2_low	liver_H0012_1_low	liver_H0012_2_low	liver_H0021_1_none	liver_H0021_2_none
liver_H0007_1_high	1.0000	0.9944	0.9861	0.9859	0.9892	0.9891	0.9910	0.9905	0.9878	0.9871	0.9851	0.9858	0.9895	0.9914	0.9882	0.9878	0.9889	0.9889
liver_H0007_2_high	0.9944	1.0000	0.9893	0.9886	0.9925	0.9924	0.9921	0.9917	0.9908	0.9910	0.9866	0.9865	0.9901	0.9923	0.9918	0.9910	0.9922	0.9930
liver_H0008_1_high	0.9861	0.9893	1.0000	0.9965	0.9911	0.9912	0.9881	0.9878	0.9924	0.9926	0.9855	0.9847	0.9831	0.9864	0.9911	0.9911	0.9897	0.9908
liver_H0008_2_high	0.9859	0.9886	0.9965	1.0000	0.9905	0.9904	0.9875	0.9868	0.9915	0.9922	0.9842	0.9840	0.9832	0.9857	0.9910	0.9910	0.9891	0.9902
liver_H0011_1_high	0.9892	0.9925	0.9911	0.9905	1.0000	0.9979	0.9911	0.9910	0.9922	0.9920	0.9862	0.9860	0.9863	0.9891	0.9928	0.9915	0.9918	0.9927
liver_H0011_2_high	0.9891	0.9924	0.9912	0.9904	0.9979	1.0000	0.9910	0.9908	0.9922	0.9922	0.9860	0.9856	0.9859	0.9886	0.9931	0.9919	0.9917	0.9927
liver_H0018_1_high	0.9910	0.9921	0.9881	0.9875	0.9911	0.9910	1.0000	0.9960	0.9916	0.9909	0.9896	0.9895	0.9919	0.9927	0.9910	0.9905	0.9915	0.9917
liver_H0018_2_high	0.9905	0.9917	0.9878	0.9868	0.9910	0.9908	0.9960	1.0000	0.9912	0.9910	0.9892	0.9891	0.9900	0.9916	0.9905	0.9905	0.9902	0.9907
liver_H0022_1_high	0.9878	0.9908	0.9924	0.9915	0.9922	0.9922	0.9916	0.9912	1.0000	0.9979	0.9871	0.9866	0.9861	0.9887	0.9921	0.9920	0.9928	0.9939
liver_H0022_2_high	0.9871	0.9910	0.9926	0.9922	0.9920	0.9922	0.9909	0.9910	0.9979	1.0000	0.9869	0.9861	0.9855	0.9887	0.9924	0.9925	0.9928	0.9939
liver_H0004_1_low	0.9851	0.9866	0.9855	0.9842	0.9862	0.9860	0.9896	0.9892	0.9871	0.9869	1.0000	0.9974	0.9889	0.9908	0.9834	0.9830	0.9867	0.9867
liver_H0004_2_low	0.9858	0.9865	0.9847	0.9840	0.9860	0.9856	0.9895	0.9891	0.9866	0.9861	0.9974	1.0000	0.9890	0.9910	0.9830	0.9831	0.9864	0.9858
liver_H0009_1_low	0.9895	0.9901	0.9831	0.9832	0.9863	0.9859	0.9919	0.9900	0.9861	0.9855	0.9889	0.9890	1.0000	0.9961	0.9854	0.9854	0.9877	0.9876
liver_H0009_2_low	0.9914	0.9923	0.9864	0.9857	0.9891	0.9886	0.9927	0.9916	0.9887	0.9887	0.9908	0.9910	0.9961	1.0000	0.9885	0.9885	0.9904	0.9905
liver_H0012_1_low	0.9882	0.9918	0.9911	0.9910	0.9928	0.9931	0.9910	0.9905	0.9921	0.9924	0.9834	0.9830	0.9854	0.9885	1.0000	0.9971	0.9920	0.9931
liver_H0012_2_low	0.9878	0.9910	0.9911	0.9910	0.9915	0.9919	0.9905	0.9905	0.9920	0.9925	0.9830	0.9831	0.9854	0.9885	0.9971	1.0000	0.9915	0.9924
liver_H0021_1_none	0.9889	0.9922	0.9897	0.9891	0.9918	0.9917	0.9915	0.9902	0.9928	0.9928	0.9867	0.9864	0.9877	0.9904	0.9920	0.9915	1.0000	0.9968
liver_H0021_2_none	0.9889	0.9930	0.9908	0.9902	0.9927	0.9927	0.9917	0.9907	0.9939	0.9939	0.9867	0.9858	0.9876	0.9905	0.9931	0.9924	0.9968	1.0000

Table 3. Pearson correlation coefficients of transcriptome data of all samples versus each other.

Data record 3

ELISA measurements have been deposited at figshare (<http://www.figshare.com>) (Data Citation 3).

Technical Validation

Transcriptomic data

Microarray data passed the proprietary Illumina quality controls. All samples were investigated in duplicates. Fig. 2b shows that—as would be expected—the duplicates cluster together demonstrating the validity of experiments in terms of whole-genome gene expression. The Pearson correlation coefficients of all samples versus each other were calculated with the intention to detect outliers. However, all correlation coefficients were greater than 0.98 and all correlation coefficients of duplicates even greater than 0.99 so that all samples passed this quality check (Table 3). Genes with significant differential gene expression were selected for validation via RT-PCR experiments (Fig. 2c). Genes were termed differentially expressed if the multiple-testing-corrected limma²⁸ P -value was less than 0.05, the ratio was less than 0.75 or greater than 1.33 and the gene was expressed (detection P -value less than 0.05) in at least one of both cases. Furthermore, we analysed clusters of genes differentially expressed in high versus low steatotic livers together with genes found in literature^{19,23} and genome-wide association studies³ (Fig. 2d). This analysis confirms high similarity between duplicates and clustering—to some extent but not fully—according to steatosis grade. Fig. 3a shows a plot of the first two components of the Principal Component Analysis (PCA) of the microarray data.

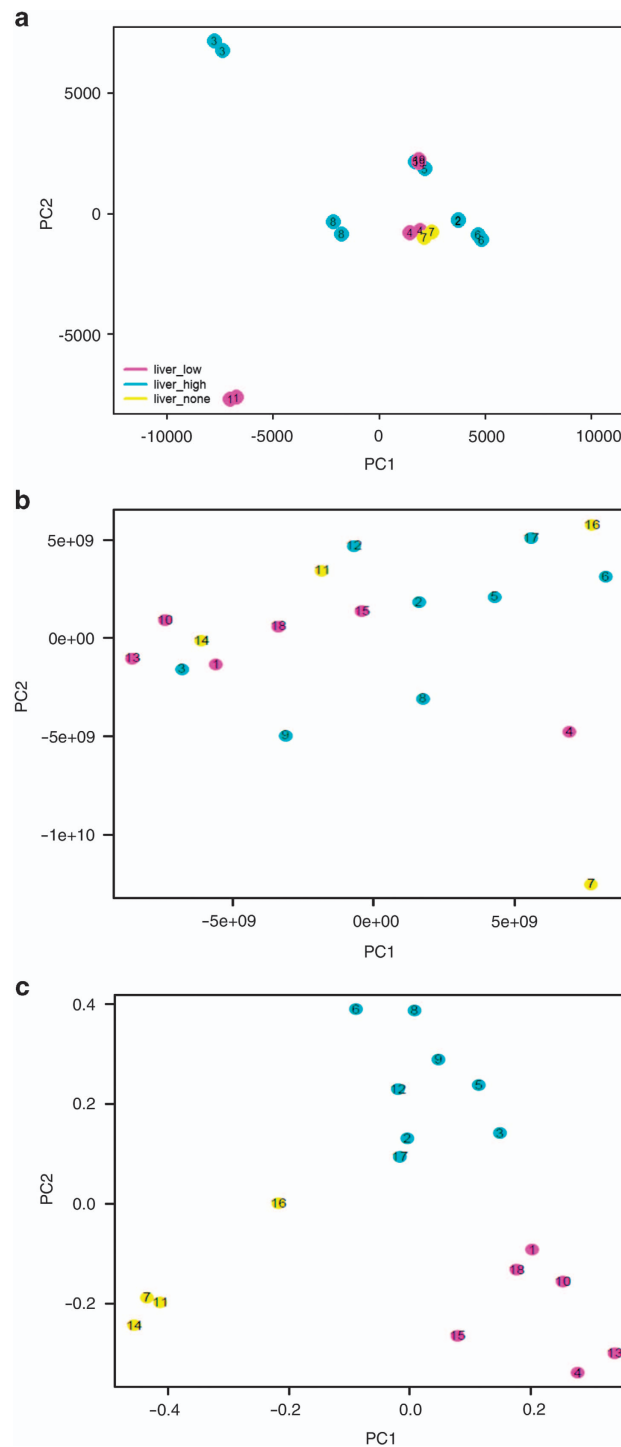


Figure 3. Transcriptomic and metabolomic profiles. (a–c) Transcriptomics and metabolomics PCA plots. Distinct colours are used to aid visualizing patients with distinct levels of steatosis: yellow, patients with < 5% no steatosis; magenta, patients with 5–33%, low level steatosis; cyan, patients with high steatosis >33%, high steatosis. (a) Unsupervised PCA plot for 18 liver biopsies, Illumina microarray data. (b) Unsupervised PCA plot for 18 plasma samples, metabolomics data. (c) Supervised discrimination analysis (pls/ca: partial least squares/canonical analysis) of metabolites in patient plasma samples. The correspondence between numbers in the plot and sample names in Table 1 is: 1 = H0004, 2 = H0007, 3 = H0008, 4 = H0009, 5 = H0011, 6 = H0018, 7 = H0021, 8 = H0022, 9 = H0024, 10 = H0025, 11 = H0026, 12 = H0027, 13 = H0028, 14 = H0029, 15 = H0030, 16 = H0031, 17 = H0033, 18 = H0034, 19 = H0012.

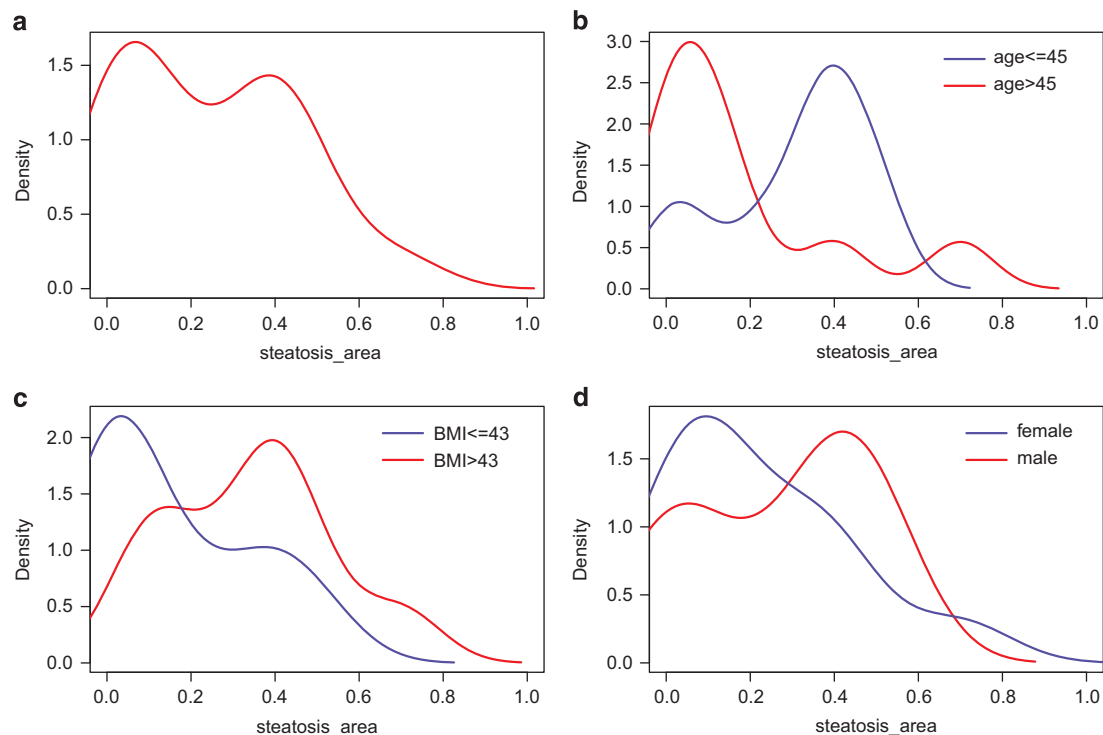


Figure 4. Distribution plots of percentage parenchymal involvement in steatosis. (a) all patients. (b) Kernel density plot of patients above/below median age (median = 45). (c) Kernel density plot of patients above/below median BMI (median = 43). (d) Kernel density plot of male/female patients.

ELISA-based assay for biomarkers

Samples below and above quantification limit as well as samples with coefficient of variation (cv) greater than 20% were marked in the measurements table (Data Citation 3). Independent validation of the ELISA-based measurements was checked by visual inspection of plots comparing disease states.

Metabolomics

Assignment of all metabolites were done manually, signals were assigned on template one-dimensional NMR profiles by using matching routines of AMIX 7.3.2 (Bruker BioSpin) in combination with the BBIREFCODE Version 2-0-0 reference database and published literature when available. Additional confirmation was done using data provided in our lab -database containing spectra of standard pure compounds. To assess which metabolites (i.e., NMR peaks) were significantly different between different sets a univariate paired Wilcoxon test was used. A P -value ≤ 0.05 was considered statistically significant (P -value not corrected for multiple testing).

Robust validation of statistical analysis results was done using a cross-validation technique.

The accuracy of the classification was assessed by means of a single cross-validation scheme. The original data set was split into a training set (80% of the samples) and a test set (20% of the samples) prior to any step of statistical analysis. The number of PLS components was chosen on the basis of a 5-fold cross validation performed on the training set only, and the best model was used to predict the samples in the test set. The whole procedure was repeated 200 times with a Monte Carlo cross validation scheme, and the results averaged.

Figure 3b shows a plot of unsupervised discrimination analysis and Fig. 3c shows separation of steatosis grades in a plot of supervised discrimination analysis (pls/ca: partial least squares/canonical analysis) of metabolites in patient plasma samples. The clustering of Fig. 3c results from a supervised PLS/CA based only on the metabolomic NMR profiles. The algorithm takes into account the supervised information relative to the 3 steatosis groups.

Distribution plots

Figure 4a shows the distribution of percentage parenchymal involvement in steatotic patients derived from Table 1. The percentage is converted to a scale from zero to one and plotted with the kernel density function from the R statistical package. Fig. 4b–d display distributions separated into groups of age above/below median (median = 45), body mass index (BMI) above/below median (median = 43) and gender. Fig. 4d would suggest a slight tendency for more severe steatosis in males. A similar trend has

been reported in a NAFLD study on Australian adolescents where 3.1% of males and only 2.2% of females had moderate to severe steatosis while 7.0% of males and 14.1% of females had mild steatosis²⁹.

Usage Notes

All patients in this study underwent bariatric surgery. This should be taken into account when generalizing results although these are typical cases of morbid obesity which is connected to the metabolic syndrome including NAFLD. The sample size of these datasets—in particular the transcriptomics dataset—poses certain limits onto its usage. Due to its small size it will not enable rigorous analysis of gender effects. Therefore it would likely need to be combined with other data sources, such as data from Moylan *et al.*¹⁸ and Du Plessis *et al.*²³.

References

- Lazo, M. & Clark, J. M. The epidemiology of nonalcoholic fatty liver disease: a global perspective. *Semin. Liver Dis.* **28**, 339–350 (2008).
- Petersen, K. F. *et al.* Apolipoprotein C3 Gene Variants in Nonalcoholic Fatty Liver Disease. *N. Engl. J. Med.* **362**, 1082–1089 (2010).
- Speliotes, E. K. *et al.* Genome-Wide Association Analysis Identifies Variants Associated with Nonalcoholic Fatty Liver Disease That Have Distinct Effects on Metabolic Traits. *PLoS Genet* **7**, e1001324 (2011).
- Samocho-Bonet, D. *et al.* Overfeeding Reduces Insulin Sensitivity and Increases Oxidative Stress, without Altering Markers of Mitochondrial Content and Function in Humans. *PLoS ONE* **7**, e36320 (2012).
- Ratziu, V., Bellentani, S., Cortez-Pinto, H., Day, C. & Marchesini, G. A position statement on NAFLD/NASH based on the EASL 2009 special conference. *J. Hepatol.* **53**, 372–384 (2010).
- Farese, R. V. Jr., Zechner, R., Newgard, C. B. & Walther, T. C. The Problem of Establishing Relationships between Hepatic Steatosis and Hepatic Insulin Resistance. *Cell Metab.* **15**, 570–573 (2012).
- Samuel, V. T. *et al.* Mechanism of Hepatic Insulin Resistance in Non-alcoholic Fatty Liver Disease. *J. Biol. Chem.* **279**, 32345–32353 (2004).
- Rajpathak, S. N. *et al.* Insulin-like growth factor axis and risk of type 2 diabetes in women. *Diabetes* **61**, 2248–2254 (2012).
- Völzke, H. *et al.* Association between hepatic steatosis and serum IGF1 and IGFBP-3 levels in a population-based sample. *Eur. J. Endocrinol.* **161**, 705–713 (2009).
- Aleem, E., Nehrbass, D., Klimek, F., Mayer, D. & Bannasch, P. Upregulation of the insulin receptor and type I insulin-like growth factor receptor are early events in hepatocarcinogenesis. *Toxicol. Pathol.* **39**, 524–543 (2011).
- Baffy, G., Brunt, E. M. & Caldwell, S. H. Hepatocellular carcinoma in non-alcoholic fatty liver disease: An emerging menace. *J. Hepatol.* **56**, 1384–1391 (2012).
- Marra, F., Gastaldelli, A., Svegliati Baroni, G., Tell, G. & Tiribelli, C. Molecular basis and mechanisms of progression of non-alcoholic steatohepatitis. *Trends Mol. Med.* **14**, 72–81 (2008).
- Angulo, P. Nonalcoholic fatty liver disease. *N. Engl. J. Med.* **346**, 1221–1231 (2002).
- Jozefczuk, J. *et al.* A Systems Biology Approach to Deciphering the Etiology of Steatosis Employing Patient-Derived Dermal Fibroblasts and iPS Cells. *Front. Physiol* **3**, 339 (2012).
- Houstis, N., Rosen, E. D. & Lander, E. S. Reactive oxygen species have a causal role in multiple forms of insulin resistance. *Nature* **440**, 944–948 (2006).
- Findeisen, H. M. *et al.* Glutathione Depletion Prevents Diet-Induced Obesity and Enhances Insulin Sensitivity. *Obesity* **19**, 2429–2432 (2011).
- Hayes, J. D., Flanagan, J. U. & Jowsey, I. R. Glutathione Transferases. *Annu. Rev. Pharmacol. Toxicol.* **45**, 51–88 (2005).
- Moylan, C. A. *et al.* Hepatic gene expression profiles differentiate presymptomatic patients with mild versus severe nonalcoholic fatty liver disease. *Hepatol. Baltim. Md* **59**, 471–482 (2014).
- Kahali, B. *et al.* TM6SF2: catch-22 in the fight against nonalcoholic fatty liver disease and cardiovascular disease? *Gastroenterology* **148**, 679–684 (2015).
- Kozlitina, J. *et al.* Exome-wide association study identifies a TM6SF2 variant that confers susceptibility to nonalcoholic fatty liver disease. *Nat. Genet.* **46**, 352–356 (2014).
- Shungin, D. *et al.* New genetic loci link adipose and insulin biology to body fat distribution. *Nature* **518**, 187–196 (2015).
- Feldstein, A. E. *et al.* Cytokeratin-18 fragment levels as noninvasive biomarkers for nonalcoholic steatohepatitis: A multicenter validation study. *Hepatology* **50**, 1072–1078 (2009).
- Du Plessis, J. *et al.* Association of Adipose Tissue Inflammation With Histologic Severity of Nonalcoholic Fatty Liver Disease. *Gastroenterology* **149**, 635–648, e14 (2015).
- Matthews, D. R. *et al.* Homeostasis model assessment: insulin resistance and β -cell function from fasting plasma glucose and insulin concentrations in man. *Diabetologia* **28**, 412–419 (1985).
- Kleiner, D. E. *et al.* Design and validation of a histological scoring system for nonalcoholic fatty liver disease. *Hepatology* **41**, 1313–1321 (2005).
- Gentleman, R. C. *et al.* Bioconductor: open software development for computational biology and bioinformatics. *Genome Biol.* **5**, R80 (2004).
- Du, P., Kibbe, W. A. & Lin, S. M. lumi: a pipeline for processing Illumina microarray. *Bioinformatics* **24**, 1547–1548 (2008).
- Smyth, G. K. Linear Models and Empirical Bayes Methods for Assessing Differential Expression in Microarray Experiments. *Stat. Appl. Genet. Mol. Biol.* **3** doi:10.2202/1544-6115.1027 (2004).
- Ayonrinde, O. T. *et al.* Gender-specific differences in adipose distribution and adipocytokines influence adolescent nonalcoholic fatty liver disease. *Hepatol. Baltim. Md* **53**, 800–809 (2011).

Data Citations

- Wruck, W. *et al.* *NCBI GEO* GSE46300 (2014).
- Wruck, W. *et al.* *MetaboLights* MTBLS174 (2015).
- Wruck, W. *et al.* *Figshare* <http://dx.doi.org/10.6084/m9.figshare.1333564> (2015).

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Author Contributions

W.W. performed transcriptomics analyses, UK and D.B. performed RPPA analyses, E.G. and P.T. NMR analyses, S.R. and H.W. flux analyses (data not included here). K.K. and K.Z. were responsible for patient recruitment and RNA isolation from patient-derived liver biopsies. J.J. and K.W. performed the RT-PCR experiments. A.D. and C.W. performed pathway analysis and built systems biological models (data not included here) and V.P. performed flux coupling analysis (data not included here). W.W. was responsible for data management. W.W., A.D., S.R., H.W., C.W. and J.A. wrote the manuscript. H.L. and J.A. conceived the concept. J.A. coordinated the work.

Additional Information

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