INTRODUCTION

Sudden infant death syndrome (SIDS) is the leading cause of death in infants older than 1 week of age\(^1\) and its etiology is still largely obscure\(^2,3\). The triple risk theory\(^4\) involves the intersection of (i) a vulnerable infant, (ii) a critical developmental period, and (iii) exogenous stressors as cause for SIDS. The vulnerability is attributed mainly to an impaired autonomic nervous system, most notably concerning the respiratory regulation and/or arousal reaction\(^5\). Serotonergic neurons constitute a major modulating system that controls the respiratory activity and these functions are adversely affected as demonstrated by numerous reports on multiple abnormalities in the serotonergic system of the brain stem\(^6-8\). Furthermore, the influence of genetic variants is discussed: several studies have been published on gene variants of relevance for the serotonergic system and other parts of the autonomous nervous system\(^9,10\).

The most prominent gene in that respect is the serotonin transporter (5-HTT) gene that is responsible for the reuptake of released serotonin (5-HT) and thus regulates the level of 5-HT in the synaptic cleft\(^13\). A promoter polymorphism of the 5-HTT gene (5-HTTLPR) alters the transcription rate of the gene: a promoter with the short “S” allele (14 copies of a 22–23 bp repeat region) is less active in vitro compared to the longer “L” allele (16 copies)\(^14,15\). After an initial description of an increased frequency of L-alleles in a small Japanese SIDS group\(^16\), Weese-Meyer corroborated this finding in Caucasians and Afro-Americans and found another 5-HTT gene polymorphism, a variable number of tandem repeat (VNTR) polymorphism in intron 2, observing an accumulation of the rare allele \(9\) was found to be associated with SIDS\(^17,18\). While the latter finding could not be confirmed by other studies, up to now the 5-HTTLPR was found to be associated in SIDS in an Italian and a Norwegian sample\(^19,20\) but not in a Swiss sample\(^21\). Even more so, Paterson et al.\(^22\) failed to demonstrate such an association in a sample set from the San Diego Medical Examiner’s Office (the “San Diego dataset”). The authors proposed that, at best, the 5-HTTLPR genotype is only a weak determinant of the risk to succumb to SIDS\(^23\). They further propose that studies of appropriate size and ethnically matched case and control datasets need to be performed.
performed before finally evaluating the role of 5-HTT gene variation in SIDS.

However, the activity of a gene is not determined by sequence variance alone. Epigenetics play an important role in developmental concerns and differentiation as well as various diseases. Furthermore, for the 5-HTT gene an epigenetic effect was demonstrated for many, mostly psychiatric conditions.

CpG sites are regions of DNA where a cytosine nucleotide is followed by a guanine nucleotide. Genomic regions that are characterized by a high frequency of CpGs are referred to as CpG islands and are common in promoter regions, especially close to the transcriptional start site. The degree of methylation of these DNA stretches has an important regulatory function. Even more so, it is known that age has a great impact on DNA methylation insofar as it increases with age. The infant has to adapt to the new environment after birth and grows rapidly during the first year with numerous metabolic challenges. It can be assumed that these adaptation processes are at least in part mediated by epigenetic changes. Therefore, the critical developmental period postulated by the triple risk model might well be the consequence of an inadequate epigenetic regulation of genes. Thus, we hypothesized, that the 5-HTT gene might have an impact on the etiology of SIDS not so much by genetic variants but by the degree of methylation.

To test this hypothesis we genotyped the 5-HTTLPR, the intron 2 VNTR polymorphism, and a rare gene variant which leads to gain of function (Ile428Val) in the currently largest sample tested. Also, we used bisulfite sequencing to identify the methylation status of the immediate proximal promoter of the serotonin transporter gene to review its suitability as a biomarker.

**METHODS**

**Typing of 5-HTT polymorphisms**

**Subjects.** The subjects used for this part of the study were 274 Caucasian infants from Germany whose death occurring during sleep remained unexplained after a thorough investigation, including a complete autopsy, the review of the circumstances of death, and the clinical history, as demanded by Krous et al. for establishing the diagnosis "SIDS". 175 of the infants were males and 99 females. The age range was between the 5 and 360 days (median 110 days). For the Ile428Val study of those cases, 65 cases were selected (age range 28–292 days, median 122 days). For most cases, information concerning the age at death and the date of death was available; for a minority, only information concerning the position in infants at death were available (40 SIDS victims were found in a prone position, for the remaining either no information was documented or they were found in another position). Only for some of the cases information regarding maturity at birth was available, wherefore postnatal age rather than postconceptual age was used for further analysis.

The controls consisted of 264 healthy young adult Caucasian persons who had survived the critical time span of the first year without SIDS (110 females, 154 males). Forty-four of these controls were included in the Ile428Val study. Due to anonymization, no data on age are available. The study was approved by the local ethics committee.

For the methylation study, 33 SIDS cases (age range 18–305 days, median 99 days, 21 males, 14 females) and 14 controls (age range 5–608 days, median 111 days, 8 males, 6 females) were included. As (due to the rarity of other causes of death prevalence of SIDS as cause of death) control cases are rare, we decided to include also children for up to the end of the second year. The control cases died from heart defect (4), shaken impact syndrome (4), infectious disease (4), ischemia of the colon (1), and an inborn metabolic defect (1). Three types of tissue were investigated (blood, pons cerebri, and medulla oblongata), but (due to missing samples) for pons only 24 cases and 7 controls and for the medulla only 21 cases and 6 controls could be included.

DNA analysis. Genomic DNA was extracted from 0.5 mL aliquots of blood samples or tissue samples, as described previously. Each 25 μL assay contained 2.5 μL 10× PCR buffer (including 15 mM MgCl2), 0.5 μL dNTP mix (300 μM final concentration, 0.3 μM of each primer, and 0.2 μL Taq Gold Polymerase (Life Technologies, Darmstadt, Germany). The 5-HTTLPR polymorphism was genotyped using primers as described (forward: 5’TCC TCC GCC TTG GGG CTT CC-3’ and reverse: 5’TGG GTG TTT GAG GGG AGA TCC TG-3’). After an initial denaturation step (95 °C for 15 min), thermocycling was performed for 37 cycles of 94 °C/30 s, 65 °C/60 s and 72 °C/60 s, followed by a final extension period of 72 °C for 10 min using a FlexCycler (Analytik Jena, Jena, Germany) resulting in fragments of 484 bp (S) and 528 bp (L). These fragments were separated and detected on 2% agarose gels in the presence of ethidium bromide and visualized under UV light.

For the intron 2 VNTR polymorphism, primer sequences used were forward: 5’TGG AAC CCC CAC CCT CCT CC-3’ and reverse: 5’CAT CCT CCC ACA GCA TTC TAC C-3’ (95 °C for 11 min), 37 cycles of 94 °C/30 s, 58 °C/30 s, and 72 °C/90 s, followed by a final extension period of 72 °C for 10 min. After restriction with BclI for 2.5 h at 37 °C (New England Biolabs, Frankfurt, Germany), samples were typed on a QiAxlcel electrophoresis unit using the QiAxlcel DNA Screening Kit (Qiagen, Hilden, Germany) as previously described. Fragment lengths were 365 bp (unrestricted) and 288/77 bp (restricted).

Methylation analyses

For each individual, up to three tissue samples (blood, medulla oblongata, and pons cerebri) were analyzed. DNA was extracted from tissue saved during the autopsies using the QIAamp DNA Mini Kit (Qiagen, Hilden, Germany) for blood samples on filter paper and the NucleoMag Blood 200 μL Kit (Macherey-Nagel, Düren, Germany) for deep frozen tissue samples. DNA concentration was determined using the Qubit dsDNA HS Assay Kit (Invitrogen, Carlsbad, CA).

DNA samples were bisulfite-converted and purified using the Epitect 96 Bisulfite Kit (Qiagen, Hilden, Germany) as previously described. Primers to amplify the region 28563061–28563299 of the internal promoter region were purchased from Metabion international, Planegg/Steinkirchen, Germany (812_5HTTE-Fw-N (GAATTGTAGTTGGTTAATAAAATTA TT) and reverse: 5’GGT TTT GAG GGG AGA TCC TG-3’). The forward primer was labeled using the EpiTect 96 PCR Kit (Qiagen, Hilden, Germany) as previously described. PCR targeting SLCA46 was performed using an extension temperature of 58.5 °C for 35 cycles and the purified product sequenced on a Genetic Analyser 3500 XL (Applied Biosystems, Foster City, CA) after applying POP3.1 ddNTP chemistry for the sequencing PCR (ABI Life Technologies, Grand Island).

**Statistical analyses**

Exact tests for deviation from Hardy–Weinberg equilibrium were performed using the Genepop software (http://wbiomed.curtin.edu.au/genepop/). The algorithm of Roff and Bentzen was applied to test differences in the allele frequencies between controls and SIDS cases in male and female individuals and genotype frequencies in female individuals using the RxCaret software (George Carmody, Carleton University, Canada). Statistical significance was calculated by performing 10,000 simulations on the overall observed two-way contingency tables. The null hypothesis was rejected at a level of >0.05.

Analysis of CpG-specific methylation percentage was performed using the EpiGenetic Sequencing Methylation (ESME) analysis software and the methylation rate (%) of each CpG site within the amplified region was estimated by the ratio between normalized peak values of cytosine (C) and thymine (T) as described.

**RESULTS**

All investigated loci were in Hardy–Weinberg equilibrium Allele frequencies and genotype frequencies of the 5-HTTLPR polymorphism in SIDS and control subjects are summarized in Table 1 (left). For 273 SIDS cases and 277 controls, results were
and the frequency of this allele showed a non-significant trend towards association with SIDS. Pooling data for Caucasoid populations from all previously published studies resulted in a SIDS group of 438 cases vs. a control group of 485 individuals. As expected, allele 9 was the most frequent in SIDS compared to controls, although this difference did not reach statistical significance. As in previous studies, three different alleles with 9, 11, and 12 repeats were detected in a total of 504 individuals. As expected, allele 9 was the rarest of these alleles, and we observed it almost twice as frequently in SIDS compared to controls, although this difference was not significant.

For the intron 2 VNTR polymorphism, stratifying our data according to sex, age, season, and position at death demonstrated a significant association for those SIDS cases that died at 151–365 days ($p: 0.014$) and those that died during autumn ($p: 0.048$), in both cases showing a predominance of alleles 9 and 10 compared to allele 12 (Table 3).

When genotyping the Ile425Val SNP, we failed to demonstrate a significant trend for both SIDS cases and controls. For the intron 2 VNTR polymorphism, stratifying our data according to sex, age, season, and position at death demonstrated a significant association for those SIDS cases that died at 151–365 days ($p: 0.048$), in both cases showing a predominance of alleles 9 and 10 compared to allele 12 (Table 3).

At the right side, the results for the pooled data are compared for allele 9 vs. all other alleles combined. n.s. not significant.

DISCUSSION

Despite the fact that no Mendelian mode of inheritance was demonstrated for SIDS, it is assumed that multiple genetic influences contribute to the risk to succumb to this syndrome. Thus, various gene variants have been investigated in SIDS, among which the 5-HTTLPR polymorphism was the first polymorphism of relevance for the serotonergic system to be associated with SIDS. However, as some studies failed to reproduce this effect and others obtained. In accordance with the findings of Paterson et al., the frequency of the L-allele in Caucasian controls was well below 50% and the frequency of this allele showed a non-significant trend towards association with SIDS. Pooling data for Caucasoid populations from all previously published studies resulted in a SIDS group of 438 cases vs. a control group of 485 individuals. For these pooled groups, a strong association of allele L with SIDS was demonstrated (Table 1, right).

Results for the intron 2 VNTR are displayed in Table 3. For 240 cases and 264 controls, results were obtained. As in previous studies, three different alleles with 9, 11, and 12 repeats were detected in a total of 504 individuals. As expected, allele 9 was the rarest of these alleles, and we observed it almost twice as frequently in SIDS compared to controls, although this difference was not significant.

After stratifying our data for the 5-HTTLPR polymorphism, we could demonstrate a significant association with allele L for those SIDS cases that died at 46–150 days ($p: 0.045$) and those that died in a prone position ($p: 0.011$) (Table 3, left).

For the intron 2 VNTR polymorphism, we could demonstrate a significant association with allele L for those SIDS cases that died at 46–150 days ($p: 0.045$) and those that died in a prone position ($p: 0.011$) (Table 3, left).

| Allele | SIDS | Controls | Allele | SIDS | Controls |
|--------|------|----------|--------|------|----------|
| $S$    | 227  | 41.58%   | $S$    | 370  | 42.20%   |
| $L$    | 319  | 58.42%   | $L$    | 506  | 57.80%   |
| $p$    | n.s. |          | $p$    | 0.001| $0.0010$ |
| S/S    | 51   | 18.68%   | S/S    | 83   | 18.90%   |
| L/S    | 125  | 45.79%   | L/S    | 204  | 46.60%   |
| L/L    | 97   | 35.53%   | L/L    | 151  | 34.50%   |
| Total  | 273  | 100%     | Total  | 438  | 100%     |
| $p$    | n.s. |          | $p$    | 0.014| $0.0037$ |

n.s. not significant.
showed only a weak association, it is still unclear if, and if so to what extent, this polymorphism is indeed associated with SIDS. In that respect, the 5-HTTLPR polymorphism is similar to the MAOA promoter polymorphism, another polymorphism of relevance for the serotonergic system that was associated with SIDS, for which also widely discrepant studies were published. However, a recently published meta-analysis for this polymorphism results in an association with SIDS. In that respect, the 5-HTTLPR polymorphism is similar to the "5-HTT gene in association with SIDS. In fact, to the best of our knowledge, to date only one methylation study was published on SIDS cases, targeting a CpG island located upstream of the promoter of the growth factor independent 1 (GFI1) gene depending on the fact whether the parents of the deceased infant were smokers or not.

Methylation adapts the genome to changing requirements due to diseases, chronic intoxications, ageing, and even traumatic life events. The first year of life is a time of constant adaptation to changing needs. On these grounds, one of the theoretic approaches to SIDS is that the infant is maladapted to the current environmental and biological conditions. Differences in DNA methylation might very well explain this maladaptation. It is therefore somewhat surprising that the epigenetics of SIDS have been largely neglected up to now. In our study all children, controls, as well as SIDS cases lacked any methylation in the internal promoter region. Although the lack of methylation reported herein does not support a potential epigenetic influence in the etiology of SIDS, we propose to extend epigenetic studies to other genes that might be of relevance for current concepts in SIDS, i.e. the triple risk hypothesis, and of which some have already been, with rather ambiguous results, been associated to SIDS on the genomic level.

Age and season at death are external factors of high importance in SIDS, with peak SIDS prevalences between the second and fourth month of life and during the cold months of the year, respectively. Also, boys are at an approximately 50% higher risk than girls. The most influential factor determining the SIDS risk, however, is a prone sleeping position. As SIDS is a multifactorial syndrome (comprising deaths from arrhythmias, metabolic, immunologic, or respiratory derailment), we speculate that the presence or absence of one of these factors could help to further categorize Sudden infant death: Infants dying in a prone position, for instance, might more often die from brain stem- and serotonin-related respiratory failure, whereas children found in a supine position might show a preponderance of other causes. For example, in an earlier study, we succeeded to demonstrate an association with the age of 46–150 days for the THO1 locus. We therefore stratified our data for the SIDS group according to these factors and compared these groups to the control group. For age, sex, and season this information was available for almost all samples. However, only for 40 deceased infants it was documented that those were found in a prone position. The subgroups were compared to the controls rather than the remaining SIDS samples.

Most interestingly, the conditions with the highest prevalence of SIDS (age 46–150 days and prone position) are those with a significant correlation to allele L of 5-HTTLPR (Table 3). This finding appears to support the notion that the subtype of SIDS appears to actually vary depending on the age and sleeping position at death.

However, it must be noted that Opdal et al. found an association between S/S genotype and prone position—not the

| Table 3. Results for 5-HTTLPR and 5-HTT intron 2 VNTR polymorphisms in SIDS cases after stratification according to age at death, sex, season, and sleeping position compared to living controls. |
|-----------------|-----------------|-----------------|-----------------|-----------------|
| S-HTTLPR        | S-HTT intron 2  |
| 5-HTT intron 2  |
| Age (days)      |                  |
| 0–45            |                  |
| 12              |                  |
| 46–150          |                  |
| 12              |                  |
| 151–365         |                  |
| 12              |                  |
| Sex             |                  |
| m               |                  |
| f               |                  |
| Season          |                  |
| Spring          |                  |
| Summer          |                  |
| Autumn          |                  |
| Winter          |                  |
| Position        |                  |
| Prone           |                  |

Bold values indicate statistical significance p < 0.05.
L-allele as in our study. This finding emphasizes that size is a crucial quality in association studies and that detailed conclusions are only possible after a sufficient number of sufficiently large studies (our study could include only 40 infants with known prone position, that of Opdal and colleagues comprised 93 infants with prone position). Future studies should focus on the position in which the child was found.

Further, we found a very weak association for intron 2 VNTR with autumn and a stronger one with death during days 151–365 of life. However, in both cases not the initially reported allele 12, but the alleles 9 and 10 did predominate. While we consider the association with autumn most likely a statistical artifact, the association of this locus with older children seems to be more plausible, and one explanation might be that both loci have different impact on the serotonergic system.

In conclusion, we succeeded to demonstrate significant associations of two polymorphisms studied herein (5-HTTLPR and intron 2 VNTR) with SIDS, albeit only after stratification according to position, age, and season at death. We argue that stratification of SIDS samples might be a valuable approach in studies concerning the effect of age, and season at death. We argue that stratification of SIDS samples might be a valuable approach in studies concerning the genetic background of SIDS, as it could enable the researcher to reduce the bias caused by the diverse etiology of SIDS. Moreover, when pooling our results with those found in the literature, we found significant associations for both polymorphisms. Meta-analytic studies are a method to increase the power of an association study but are undoubtedly limited and prone to inherent problems. Yet, our results suggest that both polymorphisms indeed contribute to the risk of dying suddenly during the first year of life, but that the risk conferred by these alleles is relatively small, especially when compared to that conferred by some external risk factors like, e.g., the prone vs. supine sleeping position.

REFERENCES

1. Krouss, H. F. et al. Sudden infant death syndrome and unclassified sudden infant deaths: a definitional and diagnostic approach. Pediatrics 114, 234–238 (2004).
2. McCormack, J. The role of genetic testing in paediatric syndromes of sudden death: state of the art and future considerations. Cardiol. Young 2(3 Suppl.), 54–65 (2009).
3. Opdal, S. H. & Rognum, T. O. The sudden infant death syndrome gene: does it exist? Pediatrics 114, 506–512 (2004).
4. Filiano, J. J. & Kinney, H. C. A perspective on neuropathologic findings in victims of the sudden infant death syndrome: the triple-risk model. Biol. Neonate 65, 194–197 (1994).
5. Kinney, H. C., Richerson, G. B., Dymecki, S. M., Dannall, R. A. & Nattie, E. E. The brainstem and serotonin in the sudden infant death syndrome. Annu. Rev. Pathol. 4, 517–550 (2009).
6. Duncan, J. R. et al. Brainstem serotonergic deficiency in sudden infant death syndrome. JAMA 303, 430–437 (2010).
7. Machaalanl, R., Say, M. & Waters, K. A. Serotonergic receptor 1A in the sudden infant death syndrome brainstem medulla and associations with clinical risk factors. Acta Neuropathol. 117, 257–265 (2009).
8. Broadbelt, K. G. et al. Brainstem deficiency of the 14-3-3 regulator of serotonin synthesis: a proteomics analysis in the sudden infant death syndrome. Mol. Cell Proteomics 11, M111 (2012).
9. Cummings, K. J. et al. Sudden infant death syndrome (SIDS) in African Americans: polymorphisms in the gene encoding the stress peptide pituitary adenylate cyclase-activating polypeptide (PACAP). Acta Paediatr. 98, 482–489 (2009).
10. Koz, J., Schürmann, P., Rothämel, T., Dörk, T. & Klintschar, M. Gene variants associated with obstructive sleep apnea (OSA) in relation to sudden infant death syndrome (SIDS). Int. J. Legal Med. https://doi.org/10.1007/s00414-020-02480-0 (2021).
11. Klintschar, M. & Heimbold, C. Association between a functional polymorphism in the MAOA gene and sudden infant death syndrome. Pediatrics 129, e756–e761 (2012).
12. Klintschar, M., Reichenpflader, B. & Saternus, K. S. A functional polymorphism in the tyrosine hydroxylase gene indicates a role of noradrenergic signaling in sudden infant death syndrome. J. Pediatr. 153, 190–193 (2008).
13. Blakely, R. D., Defelice, L. J. & Galili, A. Biogenic amine neurotransmitter transporters: just when you thought you knew them. Physiology (Bethesda) 20, 225–231 (2005).
14. Lesch, K. P. & Mosner, R. Genetically driven variation in serotonin uptake: is there a link to affective spectrum, neurodevelopmental, and neurodegenerative disorders? Biol. Psychiatry 44, 179–192 (1998).
15. van Dyck, C. H. et al. Central serotonin transporter availability measured with [123I]beta-CIT SPECT in relation to serotonin transporter genotype. Am. J. Psychiatry 161, 525–531 (2004).
16. Narita, N. et al. Serotonin transporter gene variation is a risk factor for sudden infant death syndrome in the Japanese population. Pediatrics 107, 690–692 (2001).
17. Weese-Mayer, D. E. et al. Sudden infant death syndrome: association with a promoter polymorphism of the serotonin transporter gene. Am. J. Med. Genet. 117A, 268–274 (2003).
18. Weese-Mayer, D. E. et al. Association of the serotonin transporter gene with sudden infant death syndrome: a haplotype analysis. Am. J. Med. Genet. 122A, 238–245 (2003).
19. Nonnis Marzano, F. et al. Genes regulating the serotonin metabolic pathway in the brain stem and their role in the etiopathogenesis of the sudden infant death syndrome. Genomics 91, 485–491 (2008).
20. Opdal, S. H., Vege, A. & Rognum, T. O. Serotonin transporter gene variation in sudden infant death syndrome. Acta Paediatr. 97, 861–865 (2008).
21. Haas, C., Braun, J., Bär, W. & Bartsch, C. No association of serotonin transporter gene variation with sudden infant death syndrome (SIDS) in Caucasians. Leg. Med. (Tokyo) 11, S210–S212 (2009).
22. Paterson, D. S. et al. Lack of association of the serotonin transporter polymorphism with the sudden infant death syndrome in the San Diego Dataset. Pediatr. Res. 68, 409–413 (2010).
23. Paterson, D. S. Serotonin gene variants are unlikely to play a significant role in the pathogenesis of the sudden infant death syndrome. Respir. Physiol. Neurobiol. 189, 301–314 (2013).
24. Moore, L. D., Le, T. & Fan, G. DNA methylation and its basic function. Mol. Psychiatry 38, 23–38 (2013).
25. Bakkus, J., Schaufeli, W., Claes, S., & Godderis, L. Stress, burnout and depression: a systematic review on DNA methylation mechanisms. J. Psychosom. Res. 92, 34–44 (2017).
26. Bird, A. DNA methylation patterns and epigenetic memory. Genes Dev. 16, 6–21 (2002).
27. Ozaki, N. et al. Serotonin transporter missense mutation associated with a complex neuropsychiatric phenotype. Mol. Psychiatry 8, 933–936 (2003).
28. Laer, K., Vennemann, M., Rothämél, T. & Klintschar, M. Association between polymorphisms in the P2RY1 and SSTR2 genes and sudden infant death syndrome. Int. J. Leg. Med. 127, 1087–1089 (2013).
29. Dauber, E. M. et al. Germline mutations of STR-alleles include multi-step mutations as defined by sequencing of repeat and flanking regions. Forensic Sci. Int. Genet. 6, 381–386 (2012).
30. Wendland, J. R., Martin, B. J., Kruse, M. R., Lesch, K. P. & Murphy, D. L. Simultaneous genotyping of four functional loci of human SLC6A4, with a reanalysis of 5-HTTLPR and rs25531. Mol. Psychiatry 11, 224–226 (2006).
31. Klintschar, M. & Heimbold, C. No association of SIDS with two polymorphisms in genes relevant for the noradrenergic system: COMT and DBH. Acta Paediatr. 101, 1079–1082 (2012).
32. Rhein, M. et al. DNA methylation results depend on DNA integrity role of post mortem interval. Front. Genet. 6, 182 (2015).
33. Roff, D. A. & Bentzen, P. The statistical analysis of mitochondrial DNA polymorphisms: X2 and the problem of small samples. Mol. Biol. Evol. 6, 539–545 (1989).
34. Levin, J., Schmitt, A. O., Asofani, P., Hildmann, T. & Piepenbrock, C. Quantitative DNA methylation analysis based on four-dye trace data from direct sequencing of PCR amplificates. Bioinformatics 20, 3005–3012 (2004).
35. Weese-Mayer, D. E., Ackerman, M. J., Marazita, M. L. & Berry-Kravis, E. M. Sudden infant death syndrome: review of implicated genetic factors. Am. J. Med. Genet. A 143A, 771–788 (2007).
36. Filionz, L. et al. Association of dopamine transporter and monoamine oxidase molecular polymorphisms with sudden infant death syndrome and stillbirth: new insights into the serotonin hypothesis. Neurogenetics 10, 65–72 (2009).
37. Klintschar, M., Rothämél, T. & Lær, K. Monoamine oxidase A and sudden infant death syndrome. J. Pediatr. 163, 1533 (2013).
38. Courts, C., Grabmüller, M. & Madea, B. Monoamine oxidase A gene polymorphism and the pathogenesis of sudden infant death syndrome. J. Pediatr. 163, 89–93 (2013).
39. Zhou, Q., Gong, D., Zhang, Y. & Huang, F. Association between monoamine oxidase A promoter polymorphism and the risk of sudden infant death syndrome: a meta-analysis. Int. J. Leg. Med. https://doi.org/10.1007/s00414-020-02496-6 (2021).
AUTHOR CONTRIBUTIONS
N.P., M.R., F.M.-B., K.L. and D.G.: substantial contribution to conception and design. Acquisition, analysis and interpretation of data, drafting the manuscript or revising it critically for important intellectual content, final approval. D.Q. and H.F.: substantial contribution to conception and design. Interpretation of data, drafting the manuscript or revising it critically for important intellectual content, final approval. V.P., T.R., A.A., T.A.E., and B.V.: substantial contribution to conception and design. Analysis and interpretation of data, writing the manuscript, final approval.

FUNDING
Open Access funding enabled and organized by Projekt DEAL.

COMPETING INTERESTS
The authors declare no competing interests.

ETHICS APPROVAL AND CONSENT TO PARTICIPATE
The study was approved by the local ethics committee.

ADDITIONAL INFORMATION
Supplementary information The online version contains supplementary material available at https://doi.org/10.1038/s41390-021-01773-3.

Correspondence and requests for materials should be addressed to Mathias Rhein or Michael Klintiachar.

Reprints and permission information is available at http://www.nature.com/reprints

Publisher’s note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made. The images or other third party material in this article are included in the article’s Creative Commons license, unless indicated otherwise in a credit line to the material. If material is not included in the article’s Creative Commons license and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this license, visit http://creativecommons.org/licenses/by/4.0/.

© The Author(s) 2021