Fecal microbiota diversity in survivors of adolescent/young adult Hodgkin lymphoma: a study of twins

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Background: Adolescent/young adult Hodgkin lymphoma (AYAHL) survivors report fewer exposures to infections during childhood compared with controls, and they have functional lymphocyte aberrations. The gut microbiota plays a central role in immunity.

Methods: We investigated whether fecal microbial diversity differed between 13 AYAHL survivors and their unaffected co-twin controls. Pyrosequencing of fecal bacterial 16S rRNA amplicons yielded 252943 edited reads that were assigned to species-level operational taxonomic units (OTUs) and standardised for sequencing depth by random sampling. Microbial diversity was compared within vs between twin pairs and by case–control status.

Results: The number of unique OTUs was more similar within twin pairs compared with randomly paired participants (P = 0.0004). The AYAHL cases had fewer unique OTUs compared with their co-twin controls (338 vs 369, P = 0.015); this difference was not significant (169 vs 183, P = 0.10) when restricted to abundant OTUs.

Conclusion: In this small study, AYAHL survivors appear to have a deficit of rare gut microbes. Further work is needed to determine if reduced microbial diversity is a consequence of the disease, its treatment, or a particularly hygienic environment.

Successful treatment of adolescent/young adult Hodgkin lymphoma (AYAHL) has enabled investigation of the pathogenesis and consequences of this disease. Adolescent/young adult Hodgkin lymphoma is associated with reduced exposures to infections, including small sibship size, less crowded living conditions, and higher socioeconomic status (Westergaard et al, 1997). Similarly, a previous study of AYAHL survivors suggested fewer early childhood fecal-oral exposures compared with unaffected co-twin controls (Cozen et al, 2009). Adolescent/young adult Hodgkin lymphoma is associated with suppressed T-helper 1 (Th1) immunity, a hyper-inflammatory Th2 response, and persistent immune defects (Fisher et al, 1980; Poppema, 1996; Salas et al, 2012). Colonisation, expansion, and maturation of gut microbial populations during infancy (Dominguez-Bello et al, 2010; Koenig et al, 2011) coincide with a switch from a fetal Th2-dominated to a mature Th1-dominated immune profile (Matricardi and Bonini, 2000; Martin et al, 2010). Elevated levels of IgE and Th2 cytokines in AYAHL patients, as well as deficits of cytotoxic T cells and

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natural killer cells in Hodgkin lymphoma tumours (Romagnani et al, 1980; Samoszuk, 1992; Skinnider and Mak, 2002; Poppema, 2005), suggest a failure to make this early Th2-to-Th1 switch. These observations raise the possibility that the gut microbiota, which strongly influences adaptive and innate immunity (Blumberg and Powrie, 2012; Hooper et al, 2012), may impact AYAHL. Thus, the current study employed next generation sequencing of fecal microbial DNA to test whether gut microbial diversity or taxonomy differed between AYAHL survivors and unaffected co-twin controls.

**MATERIALS AND METHODS**

Detailed Materials and methods are in Supplementary information. Briefly, subjects were AYAHL-discriminant twin pairs participating in the International Twin Study (Mack et al, 1995). Living pairs, resident in the United States or Canada, were eligible if one member had been diagnosed with histologically verified AYAHL under 50 years of age (actual age 18–44 years), if both members of the pair were alive according to a death index linkage, and if the pair was discordant for reported early childhood fecal-oral exposures as assessed by a questionnaire completed and returned by both members of the pair (n = 33 pairs) (Cozen et al, 2009). At least 1 member of 15 pairs either declined or was not able to be located, and 5 pairs were excluded for gastrointestinal illness or recent use of certain medications in at least 1 member, leaving 13 pairs. All 26 participants provided written informed consent and 1 fecal specimen. The majority of the HL survivors had nodular sclerosis subtype that was negative for Epstein Barr virus (EBV). Participants were provided with a kit that fits over the toilet seat for fecal specimen collection. Specimens were immediately frozen at −20°C, shipped overnight to University of Southern California, and frozen at −80°C after replacement of identifying information with a code number. The study was approved by the responsible Institutional Review Boards.

Following microbial cell lysis, DNA was extracted from three aliquots of each fecal specimen, pooled, and used to amplify V2 region 16S rRNA genes using primers FWD: AGAGTTTG ATCCTACGGCTCAAG and REV: TGCTGCGTCCGTAGGACT (Turnbaugh et al, 2009). The amplicons were purified, pooled, and sequenced (454 Life Sciences FLX, standard chemistry; Roche Company, Branford, CT, USA). Low-quality and chimeric sequences were removed (Wang and Wang, 1996; Caporaso et al, 2010; Edgar et al, 2011). Remaining sequences with >97% identity were defined as species-level operational taxonomical units (OTUs) (Edgar, 2010) and assigned to taxa with the Ribosomal Data Project classifier version 2.2 (Wang et al, 2007). Each taxon’s proportion was its relative abundance.

Comparisons were performed on 5520 sequences (initial analysis) or 5258 sequences (‘conservative’ analysis that discarded rare OTUs) randomly sampled from each individual. Difference in OTU alpha diversity (within a sample) within twin pairs, compared with randomly paired individuals, was tested by one-way analysis of variance (Howell, 2002). Cases and controls were compared on alpha diversity and body mass index by paired t-test (Freeman et al, 2007). Differences in microbial communities within vs between twin pairs (beta diversity) were tested with UniFrac (weighted and unweighted for relative abundance; Lozupone and Knight, 2005) and permutation tests with 1000 replications. Each time, the sum of the paired UniFrac distances across the 13 twin pairs was compared with the corresponding sum in 13 randomly paired sets of individuals. For case–control comparisons of beta diversity, the sum of UniFrac values across all pairings of the 13 cases (13 x 12/2 = 78 summands) was compared with the corresponding sum in 1000 random selections of 13 individuals.

Monte Carlo methods (Caporaso et al, 2010) provided nearly identical results (not presented). Case–control comparisons of taxa with mean relative abundance ≥0.1% (Freeman et al, 2007) were performed with paired t-tests. All statistical tests were two-sided. *P <0.05 was deemed significant.

**RESULTS**

Table 1 presents demographic characteristics of the 13 twin pairs. Cases were diagnosed an average of 22.5 years (range 10–36 years) before participation in this study. After editing to remove low quality sequences and amplification artefacts (chimeras), there were 252 943 bacterial V2-16S rRNA reads from the 26 fecal samples. These were assigned to 2513 species-level OTUs. Table 2 presents within-pair differences in alpha diversity (Shannon index and number of unique OTUs). Mean differences were significantly smaller within pairs than between pairs (Shannon index, *P* = 0.02; OTUs, *P* = 0.0004).

In the initial analysis, all measures of alpha diversity were higher in controls than in AYAHL cases (Table 3). Controls had significantly more unique OTUs (mean difference, 31), a measure that does not weight for relative abundance. Accounting for abundance, controls still had higher diversity, but the differences were attenuated and not statistically significant (Chaol difference = 41, *P* = 0.066; PD whole tree difference = 1.6, *P* = 0.051; Shannon index = 0.2, *P* = 0.27). Alpha diversity did not differ significantly by zygosity, sex, current age, age at separation, more oral exposures in early childhood, or time since AYAHL diagnosis (*P >0.19*, data not presented). Case-minus-control differences in body mass index were not significant (median 0.5 kg m⁻², mean – 0.08 kg m⁻², *P* = 0.96).

Compared with the initial analysis, the conservative analysis used 5% fewer reads per sample (5258, Supplementary Table 1) and the data set had 81% fewer OTUs (488 vs 2513). As in the initial analysis, all measures of alpha diversity in the conservative

### Table 1. Characteristics of the 13 twin pairs, discordant for Hodgkin lymphoma, whose fecal samples were used for the study

| Characteristic | Value |
|---------------|-------|
| Race | White, no. of pairs (%) 12 (92) | Non-white, no. of pairs (%) 1 (8) |
| Zygosity | MZ, no. of pairs (%) 5 (38) | DZ, no. of pairs (%) 8 (62) |
| Gender type | MM, no. of pairs (%) 3 (23) | FF, no. of pairs (%) 6 (46) | MF, no. of pairs (%) 4 (31) |
| Other characteristics | Mean age (range) at diagnosis 29 (18–44) years | Mean interval (range) between diagnosis and participation 22.5 (10–36) years | Mean no. (range) of bacterial V2-16S rRNA sequences (initial analysis) 9798 (5520–23 755) | Mean no. (range) of bacterial V2-16S rRNA sequences (conservative analysis) 9114 (5258–22 199) |

**Abbreviations:** DZ = dizygotic; F = female; M = male; MZ = monozygotic.
Fecal microbiota in Hodgkin lymphoma survivors

Table 2. Control–case differences in Shannon index and number of unique species-level operational taxonomic units (OTUs) in the fecal samples of 13 twin pairs, discordant for Hodgkin lymphoma

| Twin pairs | Zygosity | Difference in Shannon index* | Difference in number of unique OTUs* |
|------------|----------|-------------------------------|--------------------------------------|
| 1          | DZ       | 0.92                          | 71.7                                 |
| 2          | DZ       | 1.68                          | 66.3                                 |
| 3          | DZ       | –0.17                         | –5.2                                 |
| 4          | DZ       | –0.31                         | –4.1                                 |
| 5          | DZ       | 0.49                          | 37.1                                 |
| 6          | DZ       | –0.72                         | –51.5                                |
| 7          | DZ       | 0.93                          | 48.5                                 |
| 8          | DZ       | –0.12                         | 34.9                                 |
| 9          | MZ       | –0.36                         | 34.6                                 |
| 10         | MZ       | –0.09                         | –10.6                                |
| 11         | MZ       | 0.08                          | 80.7                                 |
| 12         | MZ       | 0.15                          | 67.4                                 |
| 13         | MZ       | 0.25                          | 26.8                                 |
| Mean within-pair difference** | 0.48 | 41.5 | |
| Mean between-pair difference*** | 0.75 | 77.2 | 0.02 0.0004 |

Abbreviations: DZ = dizygotic; MZ = monozygotic.
*Control value – case value.
**Mean of absolute differences within twin pairs.
***Estimated mean of the absolute difference between values from two members chosen at random from different twin sets.
****Analysis of variance to test whether values are more similar within vs between twin pairs.

Table 3. Comparisons of alpha diversity measurements between Hodgkin lymphoma cases and co-twin controls, in 13 pairs discordant for Hodgkin lymphoma

| Measurements of alpha diversity¹ | Mean (cases) | Mean (unaffected co-twins) | Mean difference (unaffected co-twins – case difference) | P-value² |
|-------------------------------|--------------|-----------------|-------------------------------------------------------|---------|
| Initial analysis              |              |                 |                                                       |         |
| No. of unique OTUs            | 338          | 369             | 31                                                    | 0.015   |
| Shannon index                 | 5.6          | 5.8             | 0.2                                                   | 0.27    |
| Chao1                         | 533          | 574             | 41                                                    | 0.066   |
| PD_whole tree                 | 21.2         | 22.8            | 1.6                                                   | 0.051   |
| Conservative analysis         |              |                 |                                                       |         |
| No. of unique OTUs            | 183          | 196             | 13                                                    | 0.10    |
| Shannon index                 | 5.2          | 5.4             | 0.2                                                   | 0.40    |
| Chao1                         | 230          | 237             | 7                                                     | 0.47    |
| PD_whole tree                 | 13.7         | 14.6            | 0.9                                                   | 0.045   |

¹OTUs, species-level operational taxonomic units; number of unique OTUs is also referred to as ‘richness’; Shannon index, which is a conservative alpha diversity estimate that adjusts for relative abundance of each OTU, is defined as (negative) the sum over OTUs of the product of the relative abundance of the OTU times the natural logarithm of the relative abundance; Chao1 is a presence/absence of alpha diversity indicator that is bias-corrected for rare taxa; PD, phylogenetic distance; PD_whole tree is an alpha diversity estimate that reflects phylogenetic divergence among OTUs present within an individual. The initial analysis included all 16S rRNA sequence reads. The conservative analysis restricted to reads with a minimum relative abundance of 0.1%.
²P-value by paired t-tests.

Cases and controls did not differ significantly in relative abundance of bacterial phylum-, class-, order-, or family-level taxa (data not shown). Restricting case–control comparisons with the 37 genera (of 108 identified) with a mean relative abundance of ≥0.1% (Supplementary Table 2), controls had a higher relative abundance of *Actinobacteria collinsella* (control mean = 0.004, case mean = 0.002, P = 0.03), which was not significant after correction for multiple tests (Miller, 1991).

Consistent with the negligible differences in relative abundance, beta diversity did not differ between the 13 cases and random groupings of 13 participants (unweighted UniFrac P = 0.07; weighted UniFrac P = 0.31). In contrast, beta diversity was significantly smaller within twin pairs compared with randomly paired participants when relative abundance was not considered (unweighted UniFrac P = 0.01), but not when relative abundance was included (weighted UniFrac P = 0.20).

**DISCUSSION**

Our initial analysis indicated that by some measurement parameters long-term survivors of AYAHL had statistically significantly fewer unique species-level OTUs in their stool than did their unaffected co-twins. These results need to be cautiously interpreted as the absolute differences were modest and other measurements of diversity, which adjust for OTU relative abundance, were not statistically significant. In our conservative analysis, restricted to the 19% most abundant OTUs, only the PD_whole tree estimate of alpha diversity was statistically significant. Deeper sequencing could shed light on rare taxa and whether they discriminate between affected and unaffected co-twins. Based on this study, we cannot determine whether lower diversity contributed to the immune perturbations that underlie AYAHL risk, or was merely a consequence of the disease and its treatment. Two small studies suggest that various chemotherapy regimens immediately reduce and alter the composition of the gut microbiota (van Vliet et al, 2009; Zwielehner et al, 2011). However, these observations were partially confounded by antibiotic use, and long-term studies are lacking. In our study, microbial diversity was assayed many years after the case twins’ diagnoses; observed differences between discordant co-twins could have been permanently induced by the disease or its treatment, or have been present before the malignancy appeared.

Twin pairs are a conservative and effective study design, not only for human genetics, but also when close matching of maternal and early life exposures is needed. Consistent with previous twin studies (Turnbaugh et al, 2009; Yatsunenko et al, 2012), fecal microbial diversity was significantly more concordant within our twin pairs than between randomly paired individuals. Likewise, concordance in diversity was similar for mono- and dizygotic pairs, suggesting that human genetic polymorphisms have little influence on the overall bacterial phylogenetic structure of the fecal microbiota (Turnbaugh et al, 2009; Yatsunenko et al, 2012). Moreover, our findings suggest that changes during adulthood related to diet, transient illnesses or treatments, are too small to modify the relatively fixed effects of twin pairing and the propensity for or consequences of obesity, inflammatory bowel disease, and, in our data, AYAHL (Turnbaugh et al, 2009; Willing et al, 2010).

Lower oral exposure to microbes in early life was associated with a 10-fold higher risk of AYAHL (Cozen et al, 2009) and with a pro-inflammatory Th2 profile in rodents. Specifically, invariant natural killer T cells, that can secrete Th2 cytokines, accumulate in germ-free mice affecting the pattern of colonisation with a gut microbiota from conventionally raised animals (Olszak et al, 2012). Neonatal rodents fail to develop a Th1 immune response on
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