The case history is presented of a woman with multiple respiratory infections and mannose binding lectin (MBL) deficiency but no evidence of bronchiectasis who developed a chronic *Burkholderia multivorans* infection. Careful microbiological assessment is needed in patients with recurrent respiratory infection and the presence of *B. multivorans* should trigger further immunological investigation including assessment of MBL status.

**CASE REPORT**

A 40 year old non-smoking woman presented to her GP in 1997 complaining of a productive cough and malaise. She was diagnosed with acute bronchitis and responded to amoxycillin (250 mg three times daily for 7 days). In August 2000 she developed a flu-like illness associated with a productive cough, wheeze, and malaise and crackles on auscultation of her chest. Her symptoms continued for 1 month. There was a transient response to 5 days of treatment with amoxycillin (250 mg three times daily) before she relapsed with a chronic cough productive of approximately 50 ml mucopurulent sputum per day. The chest radiograph at that time was normal. In 2001 the patient required nine courses of oral antibiotics for respiratory infection. Sputum cultures in 2001 were persistently positive for *Streptococcus pneumoniae* with occasional co-isolates of *Haemophilus influenzae* and *Moraxella catarrhalis*. She was referred to the Lung Defence Clinic at this hospital in 2002.

The patient had had chronic sinusitis since the age of 12 years for which she was taking beclomethasone nasal spray (50 μg per nostril twice daily) and intermittent courses of oral trimethoprim. She was diagnosed with mild asthma by her GP which was treated with Seretide 500 Accuhaler (salmeterol/fluticasone) one puff twice daily and salbutamol Diskhaler as required. In addition, she had developed epilepsy in her teens for which she took carbemazepine 400 mg twice daily. There was a strong family history of asthma. She occasionally drank alcohol and denied any illicit substance use. On direct questioning, the patient complained of persistent heartburn and acid reflux which responded well to lansoprazole 30 mg once daily.

On examination the patient had a small goitre but was clinically euthyroid. There was no finger clubbing. Her lungs were clear and the heart sounds were normal. Abdominal examination was unremarkable.

**Investigations**

Baseline blood tests including full blood count, renal and liver function, complement levels, and serum immunoglobulins were normal. Aspergillus precipitins were negative, as was an autoantibody screen. Aspergillus RAST in 2001 was negative. The chest radiograph was normal. High resolution computed tomography (HRCT) of the chest showed bronchial wall thickening in both lower lobes only. In particular, there was no evidence of bronchiectasis. Skin prick testing against a wide range of common environmental allergens including Aspergillus *fumigatus* was negative, and spirometric parameters were within the normal range. From October 2002 to July 2003 *Burkholderia multivorans* was cultured from seven of 15 sputum samples. Persistent culture of this organism prompted further immunological investigations.

The patient exhibited a poor specific antibody response to Pneumovax II, achieving antibody titres of >1 μg/ml in only five of 10 serotypes tested, and >1.3 μg/ml in only four of the 10 serotypes. Significant antibody responses were seen to serotypes 14, 18C and 19F, the main serotypes implicated in invasive disease in children, but of the five main serotypes implicated in adult disease only serotype 14 responses achieved threshold levels. Specific antibody responses to *Neisseria meningitidis* group C and *Haemophilus* group B vaccines were normal, and there was a significant antibody response to influenza A (H1N1 and H3N2) but not to influenza B. Lymphocyte surface marker analysis showed a deficit of naïve/unprimed CD4+ CD45RA+ T cells, a substantial excess of both CD4+ CD8 dull cells (equivalent to approximately 12% of CD4+ T cells), and CD28 negative T cells within both CD4+ and CD8+ T cells, similar to changes found in adult onset antibody deficiency. The CD4:CD8 ratio was normal. The patient was shown to be a heterozygote for an MBL coding region mutation (for genotyping method see Mullighan et al). MBL protein levels were not measured. The patient also displayed the CD32 (FcγRIIA) homozygous Arg131/Arg131 allotype.

CF genetics to 85% of all known mutations were negative. The CF mutation screen was carried out using the CF29 Elucigen assay which screens for delta F508 and the other 28 most common mutations in the UK population. A normal “neutrophil respiratory burst” excluded CGD.

**Management**

Initial management at this centre consisted of an admission for 10 days treatment with intravenous cefuroxime to treat *S. pneumoniae* isolated from sputum. Despite an improvement in her malaise and reduction in both cough and sputum...
production, she relapsed within a week of discharge. At this stage a Gram negative bacillus isolated from the pretreatment sputum was identified by the reference laboratory as *B. multivorans*. *B. multivorans* was grown from further sputum samples taken in the following few weeks as the patient’s symptoms deteriorated. She was readmitted to hospital for a 14 day course of intravenous tobramycin and cefazidime. Her symptoms improved and sputum samples over the next month failed to grow *B. multivorans*. Subsequent samples associated with worsening symptoms have grown *B multivorans*.

**DISCUSSION**

*Burkholderia multivorans* forms part of the *Burkholderia* complex, a group of Gram negative organisms which are commonly found in soil and water. The *Burkholderia* complex was initially described as a plant pathogen, causing sour skin disease some years ago. Initially described as a plant pathogen, causing sour skin disease, *Burkholderia multivorans* was found to be transmissible between CF patients, which is not usually the case with *B. cepacia* complex infection. This organism in our patient with no evidence of structural airway damage on HRCT scanning. It is possible that *B. multivorans* may have been present before the patient’s first visit to our centre, but was not identified as it is not routinely looked for in samples from the community. Our local laboratory has expertise in CF microbiology and is more likely to have followed up this unusual isolate.

The multiple immunodeficiencies found in this patient undoubtedly played a vital role in her susceptibility to recurrent lower respiratory tract infections. These are summarised in table 1.

Clearance of pneumococci is dependent on serotype specific antibody to capsular polysaccharide, with a bias towards IgG2 antibody production in adults. The neutrophil receptor CD32 (FcγRIIa) has two allelic forms that exhibit differential affinity for the Fc portion of IgG2. Receptors with arginine at position 131 (FcγRIIa-Arg131) bind Fc poorly, whereas the histidine variant (FcγRIIa-His131) binds avidly. If antibody is present on the bacterial surface but Fc is bound only weakly to FcγRIIa receptors, normal phagocytosis may not occur. The Arg131/Arg131 homozygous allotype for CD32 is associated with a significantly increased incidence of bacteraemic pneumococcal pneumonia. This allotype, in addition to specific antibody deficiency to *Streptococcus pneumoniae* Pneumovax II, contributed to the overall risk of pneumococcal infection in our patient.

**Mannose binding lectin (MBL)**

MBL is a serum protein involved in innate immune defence. It is part of the collectin family of proteins, including surfactant proteins A and D, with a similar function and structure. It binds carbohydrates such as mannose on the surface of microorganisms, leading to activation of the complement pathway thereby facilitating phagocytosis. Direct opsonisation also occurs via collectin receptors on the phagocyte surface.

The human MBL genes (*MBL1* and *MBL2*) have been localised to chromosome 10q11.1–21. *MBL1* is a pseudogene and *MBL2* (the functional gene) consists of four exons,

<table>
<thead>
<tr>
<th>Classification</th>
<th>Polymorphism</th>
<th>Nomenclature</th>
<th>MBL protein levels†</th>
</tr>
</thead>
<tbody>
<tr>
<td>&quot;Structural&quot; (scaling) single nucleotide polymorphisms (within exon 1)</td>
<td>Codon 52 (Arg→Cys)</td>
<td>Allele C</td>
<td>Low (100 μg/l)</td>
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<tr>
<td></td>
<td>Codon 54 (Gly→Asp)</td>
<td>Allele B</td>
<td>Low (100 μg/l)</td>
</tr>
<tr>
<td></td>
<td>Codon 57 (Gly→Glu)</td>
<td>Allele D</td>
<td>Low (250 μg/l)</td>
</tr>
<tr>
<td>Promoter region single nucleotide substitutions</td>
<td>−550 g→c (alleles H/L)</td>
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<td>High (&gt;3500 μg/l)</td>
</tr>
<tr>
<td></td>
<td>−221 c→g (alleles X/Y)</td>
<td>LYA*</td>
<td>High (&gt;2000 μg/l)</td>
</tr>
<tr>
<td></td>
<td>+4 c→t (alleles P/Q)</td>
<td>LXA*</td>
<td>Intermediate (&lt;1000 μg/l)</td>
</tr>
<tr>
<td>Compound heterozygotes (promoter polymorphisms linked to structural polymorphisms)</td>
<td>HY−52Cys</td>
<td>HYAD</td>
<td>Intermediate (1000 μg/l)</td>
</tr>
<tr>
<td></td>
<td>LY−54Asp</td>
<td>LYB</td>
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</tr>
<tr>
<td></td>
<td>LY−57Glu</td>
<td>LYC</td>
<td>Low (200 μg/l)</td>
</tr>
</tbody>
</table>

* †Nomenclature refers to promoter polymorphism on a normal structural allele (A).

Normal neutrophil activity excluded CGD. The patient had not knowingly been in contact with other patients infected with *Burkholderia* complex and had not been admitted to hospital (a known risk factor for *Burkholderia* complex infection) before identification of the first *B multivorans* isolate. The patient was exposed to multiple antibiotics, potentially leading to the emergence of infection with unusual respiratory pathogens. However, *B multivorans* has not previously been reported in patients with bronchiectasis requiring frequent courses of antibiotics. This makes “antibiotic pressure” an unlikely sole cause of persistence of this organism in our patient with no evidence of structural airway damage on HRCT scanning. It is possible that *B multivorans* may have been present before the patient’s first visit to our centre, but was not identified as it is not routinely looked for in samples from the community. Our local laboratory has expertise in CF microbiology and is more likely to have followed up this unusual isolate.

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exon 0 being non-coding. The resulting protein consists of three oligomers, each with three identical 32 kDa polypeptide chains, and is produced in the liver.

**MBL2 polymorphisms**

These are summarised in table 2. Of the structural polymorphisms described below, A denotes a normal allele whereas B, C, and D are abnormal and may be referred to as O in some laboratories. Hence, heterozygotes are often referred to as A/O genotype in the literature.

Our patient was a compound heterozygote for the structural polymorphism C/A (that is, 52 Cys) as well as a promoter polymorphism LY/HY (—500g→c substitution). Her MBL genotype is therefore HYAC and her MBL protein levels are predicted to be low at <500 μg/l (see table 2 and Garred *et al*).

**Effect of MBL2 genotype on MBL protein levels**

MBL is a functional polymer that is critically disrupted when even one of the polypeptide chains in the polymer carries an exon 1 polymorphism (table 2). Current detection methods include antibody detection using ELISA and functional assays of MBL binding. These methods select preferentially for higher order oligomerised MBL forms so they may not detect MBL antigen in MBL variant allele carriers. Garred and colleagues analysed 1183 different serum samples using an ELISA method and a new anti-MBL monoclonal antibody. They ascertained that serum obtained from A/A individuals (that is, with normal MBL levels) contained predominantly high molecular weight MBL, while serum samples from donors with a heterozygote genotype (A/O) contained both high and low molecular weight MBL. Furthermore, only high molecular weight MBL oligomers bound efficiently to mannan, enabling complement activation. Therefore, although the variant MBL genotypes give rise to higher levels of MBL than was previously thought, this variant MBL is dysfunctional. Variant MBL is also cleared from plasma more quickly than the wild type, suggesting enhanced degradation of the abnormal protein. Measuring MBL levels is not therefore as informative as determining the genotype in deciding whether or not a patient has impaired MBL function.

**MBL2 variant alleles causing reduced levels of MBL**

have been associated with an increased susceptibility of children and adults to infections. Patients with combined variable immunodeficiency (CVID) have an earlier onset of disease if MBL2 homozygosity of abnormal mannan-binding-protein alleles in patients with bronchiectasis in a non-CF patient with no HRCT suggest trigger further immunological investigation including assessment of MBL status.

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