



Draft Genome Sequence and Complete Plasmid Sequence of *Acinetobacter lwoffii* F78, an Isolate with Strong Allergy-Protective Properties

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The hygiene hypothesis states that the tremendous increase in atopic diseases correlates significantly with less contact to microbes in childhood. Here, we report the draft genome sequence of *Acinetobacter lwoffii* F78, a rural cowshed isolate with strong allergy-protective properties that contains an 8,579-bp plasmid.

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S usceptibility to asthma and allergic diseases is complex and involves genetic variants and environmental exposure (such as exposure to bacteria, viruses, and pets), alteration of our microbiome, and the large-scale manipulation of the environment over the past century. These findings led to the hygiene hypothesis, which states that the tremendous increase in atopic diseases correlates with less contact to microbes and fewer infections in childhood. A number of epidemiological studies have shown that children who grow up in a farming environment will develop less atopic disorders later in life (1–5).

The analysis of cowshed microflora from a farming environment in Bavaria (Germany) revealed one abundant bacterium, *Acinetobacter lwoffii* isolate F78, which was able to reduce allergic reactions in mice, activate mammalian cells *in vitro*, and induce a Th1-polarizing program in dendritic cells. The allergy-protective properties of this isolate were found to be imparted by its lipopolysaccharide (6–8).

Acinetobacter spp. are common in nature and widely distributed in the hospital environment and are capable of causing nosocomial infections. The genus is able to survive on moist and dry surfaces and is present in foodstuffs and on healthy human skin. In general, *Acinetobacter* spp. are considered to be nonpathogenic to healthy individuals but may cause infections in debilitated and immunocompromised people. *A. baumannii* is the most frequently isolated species from humans, whereas *A. lwoffii* belongs to the predominant species found in food. The microorganism survives desiccation for prolonged periods and grows at a wide range of temperatures (9–11).

DNA sequencing libraries were prepared using the Nextera XT kit (Illumina, USA) according to the manufacturer's instructions. Individually tagged libraries were sequenced as a part of a flow cell as 2×300 -bp paired-end reads using the Illumina MiSeq platform. A total of 12,447,167,642 sequences were produced, and the sequences from each isolate were separately assembled using CLC Genomics Workbench version 7.0.4. The location of open reading frames and the annotation of genes were done

using RAST (http://rast.nmpdr.org), and a genetic map of the resulting contigs was generated with Mauve (12, 13). The 8,579-bp plasmid was designated pAlw-F78.

Nucleotide sequence accession numbers. This whole-genome shotgun project has been deposited at DDBJ/EMBL/GenBank under the accession numbers FLLN01000001 to FLLN01000012. The versions described in this paper are the first versions.

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REFERENCES

- Daley D. 2014. The evolution of the hygiene hypothesis: the role of earlylife exposure to viruses and microbes and their relationship to asthma and allergic diseases. Curr Opin Allergy Clin Immunol 14:390–396. http:// dx.doi.org/10.1097/ACI.00000000000101.
- Schaub B, Lauener R, von Mutius E. 2006. The many faces of the hygiene hypothesis. J Allergy Clin Immunol 117:969–977. http://dx.doi.org/ 10.1016/j.jaci.2006.03.003.
- Riedler J, Braun-Fahrländer C, Eder W, Schreuer M, Waser M, Maisch S, Carr D, Schierl R, Nowak D, von Mutius E, ALEX Study Team. 2001. Exposure to farming in early life and development of asthma and allergy: a cross-sectional survey. Lancet 358:1129–1133. http://dx.doi.org/ 10.1016/S0140-6736(01)06252-3.
- Dominguez-Bello MG, Blaser MJ. 2015. Asthma: undoing millions of years of coevolution in early life? Sci Transl Med 7:307fs39. http:// dx.doi.org/10.1126/scitranslmed.aad2741.
- 5. Arrieta MC, Stiemsma LT, Dimitriu PA, Thorson L, Russell S, Yurist-Doutsch S, Kuzeljevic B, Gold MJ, Britton HM, Lefebvre DL, Subbarao P, Mandhane P, Becker A, McNagny KM, Sears MR, Kollmann T, the Child Study Investigators, Mohn WW, Turvey SE, Finlay BB, Finlay BB. 2015. Early infancy microbial and metabolic alterations affect risk of

childhood asthma. Sci Transl Med 7:307ra152. http://dx.doi.org/10.1126/ scitranslmed.aab2271.

- Debarry J, Garn H, Hanuszkiewicz A, Dickgreber N, Blümer N, von Mutius E, Bufe A, Gatermann S, Renz H, Holst O, Heine H. 2007. *Acinetobacter lwoffii* and *Lactococcus lactis* strains isolated from farm cowsheds possess strong allergy-protective properties. J Allergy Clin Immunol 119:1514–1521. http://dx.doi.org/10.1016/j.jaci.2007.03.023.
- Hanuszkiewicz A, Hübner G, Vinogradov E, Lindner B, Brade L, Brade H, Debarry J, Heine H, Holst O. 2008. Structural and immunochemical analysis of the lipopolysaccharide from *Acinetobacter lwoffii* F78 located outside *Chlamydiaceae* with a *Chlamydia*-specific lipopolysaccharide epitope. Chemistry 14:10251–10258. http://dx.doi.org/10.1002/ chem.200800958.
- Debarry J, Hanuszkiewicz A, Stein K, Holst O, Heine H. 2010. The allergy-protective properties of *Acinetobacter lwoffii* F78 are imparted by its lipopolysaccharide. Allergy 65:690–697. http://dx.doi.org/10.1111/ j.1398-9995.2009.02253.x.
- Vaneechoutte M, Nemec A, Kämpfer P, Cools P, Wauters G. 2015. Acinetobacter, Chryseobacterium, Moraxella, and other nonfermentative Gram-negative rods, p. 813–837. In Jorgensen JH, Pfaller MA, Carroll KC,

Funke G, Landry ML, Richter SS, Warnock DW (eds.), Manual of clinical microbiology, vol 1, 11th ed, vol 1. ASM Press, Washington, DC.

- Rathinavelu S, Zavros Y, Merchant JL. 2003. Acinetobacter lwoffii infection and gastritis. Microbes Infect 5:651–657. http://dx.doi.org/10.1016/ S1286-4579(03)00099-6.
- Baldeo C, Isache C, Baldeo C, Bajwa A. 2015. A case of disseminated intravascular coagulation secondary to *Acinetobacter lwoffii* and *Acinetobacter baumannii* bacteremia. IDcases 2:70–71. http://dx.doi.org/10.1016/ j.idcr.2015.05.002.
- 12. Aziz RK, Bartels D, Best AA, DeJongh M, Disz T, Edwards RA, Formsma K, Gerdes S, Glass EM, Kubal M, Meyer F, Olsen GJ, Olson R, Osterman AL, Overbeek RA, McNeil LK, Paarmann D, Paczian T, Parrello B, Pusch GD, Reich C, Stevens R, Vassieva O, Vonstein V, Wilke A, Zagnitko O. 2008. The RAST server: Rapid Annotations using Subsystems Technology. BMC Genomics 9:75. http://dx.doi.org/10.1186/ 1471-2164-9-75.
- Darling AE, Mau B, Perna NT. 2010. progressiveMauve: multiple genome alignment with gene gain, loss and rearrangement. PLoS One 5:e11147. http://dx.doi.org/10.1371/journal.pone.0011147.