Giardiasis is a neglected disease that spreads worldwide from the Arctic [1] to the Tropics [2]. It affects nearly 2% of adults and 6% to 8% of children in developed countries worldwide. Nearly 33% of people in developing countries have had giardiasis [3]. The disease is caused by a binucleated flagellated protozoan parasite *Giardia lamblia* (synonyms: *G. intestinalis* and *G. duodenalis*) that inhabits the small intestine in humans and some other mammals. *Giardia* has two morphologically distinct developmental stages, the trophozoite and the cyst. The trophozoite resides and replicates in the upper small intestine and is responsible for disease manifestations; in the lower parts of the intestine, encystation occurs, and a thick protective cyst wall is formed in this process. The cyst is the infectious, environmentally resistant stage responsible for transmission [4]. People become infected with *Giardia* by swallowing *Giardia* cysts found in contaminated food or water. Cysts are instantly infectious once they leave the host through feces. An infected person might shed 1,000,000,000–10,000,000,000 cysts daily in their feces, and this might last for several months [3]. However, swallowing as few as ten cysts might cause someone to become ill. *Giardia* is mainly passed as anthropoprosis, but zoonotic transmission also occurs. Clinical giardiasis is varied and ranges from asymptomatic passage of cysts to abdominal cramps, nausea, acute or chronic diarrhea, malabsorption, weight loss, and failure of children to thrive in both subclinical and symptomatic disease [4,5]. Pathology in giardiasis is understood to arise in several ways. These include breakdown of the epithelial barrier, defects in the epithelial brush border, increased secretion of chloride ions, and hypermotility of the intestinal smooth muscles [5]. The factors determining the variability in clinical outcome in giardiasis are still poorly understood. However, host factors (such as genotype, type of microbiota in the gut, immune status, nutritional status, and age) as well as differences in virulence and pathogenicity of *Giardia* strains are recognized as important determinants for the severity of infection [3]. Although the actual host defense mechanisms responsible for controlling *Giardia* infections are poorly understood, many studies have demonstrated the development of adaptive immune responses as well as innate mechanisms in humans and other animals. T cells and mast cells are necessary to control the infection, whereas the role of B cells in defense is contradictory [5]. *G. lamblia* has the ability to undergo extensive variation of the surface coat antigens, called variant-specific surface proteins (VSPs), which are unique, cysteine-rich zinc finger proteins. The patterns of infection in humans and animals fail to show the expected cyclical waves of increasing and decreasing numbers of parasites expressing unique VSPs. However, selection by immune-mediated processes is suggested because switching occurs at the same time that humoral responses are first detected. This process likely delays the effectiveness of the antibody response [6].

The first description of *Giardia* had been attributed to Antonie van Leeuwenhoek, who in his letter to the Royal Society on November 4, 1681, described the presence of microorganisms in his stool [7]. However, the first microscopic drawing of morphological characteristics identifying the parasite *Giardia* (Figure 1A and 1B) was provided by Vilem Dušan Lambl in 1859 (Text S1) [8], analysing the stool of a child. The similarity with the modern photographs of *Giardia* is obvious (Figure 2). He called the observed microorganisms *Cercomonas intestinalis*. In 1888, the name was changed to *Lambia intestinalis* by Raphael Anatole Émile Blanchard. In 1915, the species was renamed to *Giardia lamblia* by Charles Wardel Stiles to commemorate the work of Alfred Giard and Vilem Dušan Lambl. Because its association with disease was not constant [9], the question as to whether *Giardia* was a pathogen or a commensal was debated for many decades. In 1981, the World Health Organization (WHO) [10] added *Giardia* to the list of parasitic pathogens, but Koch’s postulates were fulfilled only in the year 1987 when Nash and colleagues demonstrated the pathogenicity of *Giardia* infections in humans by the inoculation of volunteers with trophozoites [11]. *Giardia* may be the most common pathogenic parasitic infection in humans [4]; however, no vaccine to prevent the disease in humans is available [3].

In the next part, I would like to draw attention to Vilem Dušan Lambl, the physician and scientist who first scientifically documented this parasite by microscopy drawing. Vilem Dušan Lambl (Figure 3) was born on December 5, 1824, in Letiny, near Plzen (now in the Czech Republic; in 1824, the Czech lands belonged to the Austrian Empire) [12]. He was one of eleven children; only eight reached adulthood. Around 1845, he enrolled at the University of Prague. He studied medicine but was also interested in zoology and linguistics, particularly Slavic languages and literature. He travelled extensively, especially in Bosnia, Croatia, Serbia, and Montenegro, conducting research on the southern Slavic languages, culture, fauna, and flora, and published many articles about his trips. He was a skilled draughtsman and accompanied his articles with multiple illustrations. He was also involved in the Czech patriotic and democratic movement, which later became an obstacle to his scientific career in the Austrian Empire.

**Historical Profiles and Perspectives**

**Giardia and Vilem Dušan Lambl**

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He earned his degree in medicine on January 23, 1851, and became an assistant of Professor Václav Treitz (famous for the description of the ligament of Treitz, Treitz’s hernia, and Treitz’s uremic colitis) at the Institute for Pathological Anatomy at the University of Prague. In 1856, he became associated professor (Privatdozent in German) of pathological anatomy and histology at the Medical School (University of Prague), where he taught histopathology and pathology of cancer, and worked at Josef Lochner’s children’s hospital. In 1856, he described Lambl’s excrescences (Text S2) [13], small fibrin deposits on the aortic valve. They originate as small thrombi on endocardial surfaces (where the valve margins contact) and have the potential to embolize to distant organs. He was also the first to introduce detection of bladder cancer using analysis of cells present in urine (Text S3) [14]. During his work at Josef Lochner’s children’s hospital, he described in the stool of a five-year-old girl Giardia (Figure 1), which he called Cercomonas intestinalis (Text S1) [8]. His fame attracted to his lectures young physicians arriving from the United States, Denmark, the Netherlands, Germany, and Russia. However, after the crushing of the Revolution (1848–1849) in the Austrian Empire, Felix Schwarzenberg, minister-president (1848–1852), Alexander Bach, head of internal affairs (1849–1859), and Johann Kempen, the general-gendarmerie-inspector (1849–1859), created a neoabsolutist police state. The country was infested with Secret Police, and prisons were full of political prisoners. Lambl and two of his brothers were under Secret Police surveillance, and when Lambl applied for a
professor position, he was told by the authorities that he could not become a professor in any place within the Austrian Empire [12]. Therefore, he accepted in 1860 a position at Kharkiv University (Ukraine) and in 1861 became a full professor. Ukraine was at that time under Russian rule with a regime similar to that in Austria, but at least there were no obstacles to Lambl’s scientific work. In 1871, he moved to Warsaw (in Poland, which was at that time under Russian rule) and worked there as a professor of therapy at Warsaw University and as a hospital director [12,15]. In the years 1845–1875, he published more than 100 papers [15]. During his lifetime, Lambl published in various languages (German, Czech, Russian, Polish, French, and Italian) and used different first names; therefore, it is sometimes difficult to find all of his publications. He was baptized Wilhelm (this name is also on his work about Cercomonas intestinalis (Text S1) [8]), later changed Wilhelm to the Czech version, Vilem, and added a southern Slavic name, Dusan. Some papers are published under the name Vilem Dusan Fedorovich Lambl, some articles about culture are signed “Vilem Dušan”, papers about Lambl’s excrescences (Text S2) [13] and detection of cancer cells in urine (Text S3) [14] are signed just “Dr. Lambl,” and in sources in Russian he is listed as Душан Федорович Лямбл’ [15].

He died in Warsaw on February 25, 1895. He left in his testament 20,000 Austro-Hungarian Guldens for the Fund for Support of Czech Students at the Medical Faculty of the University in Prague and at the Technical University in Prague.

Supporting Information

Text S1 Article with the first microscopic drawing of Giardia. Full text of [8]. (PDF)

Text S2 Article with the first description of Lambl’s excrescences. Full text of [13]. (PDF)

Text S3 Article with the first description of detection of bladder cancer using the analysis of cells present in urine. Full text of [14]. (PDF)

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References


Figure 3. Vilém Dušan Lambl. (A) Photo of V. D. Lambl in his younger years. The original document is stored in the Literary Archive of the Museum of Czech Literature (PNP), Prague, Czech Republic. (B) Sketch of inhabitant of Herzegovina (Portrait of Dr. Dušan Lambl as Saracen) [Hercegovec na črtach (Podobizna MUDr. Dušana Lambla jako Saracen)] by painter Jaroslav Čermák (1861). Watercolor on paper (29 × 20 cm). The original painting is in the City Gallery Prague, Czech Republic. (C) Photo of V. D. Lambl during his years in Warsaw. Original document is stored in the Literary Archive of the Museum of Czech Literature (PNP), Prague, Czech Republic. doi:10.1371/journal.pntd.0002686.g003