Chiari malformation Type I is defined as caudal displacement of the cerebellar tonsils below the level of the foramen magnum, causing compression of the brainstem and upper cervical cord. The diagnostic criteria have been debated because the absolute degree of descent below the level of the foramen magnum may be less important than the degree of compression of the cervicomedullary junction. Individuals can remain asymptomatic despite a significant descent of the tonsils, and conversely, patients can be symptomatic with little to no caudal displacement of the tonsils.10

Compression of the spinal cord, whether caused by CM-I, by a disc herniation, or by a mass lesion, has been associated with syringomyelia. The mechanism of syringomyelia formation is poorly understood, and several hypotheses have been proposed.6,8,9,11 Some have postulated that the drainage of CSF produced by the ependymal cells of the central canal is blocked, leading to a caudal accumulation of fluid. Others have argued that, in the case of CM-I, occlusion of the foramen magnum causes the intracranial CSF pulsations to be driven into the central canal or into the spinal Virchow-Robin spaces instead of its normal egress via the spinal subarachnoid space.8 Although the theory postulated by Oldfield et al. may be the most accurate,7,14 the pathogenesis may be an amalgamation of many theories, including those by Gardner,5 Williams,22 Ball and Dayan,7 Aboulker,1,2 and Greitz.6

Regardless of mechanism, it has long been recognized that restoration of CSF flow around the spinal cord is associated with a resolution of syringomyelia.21 The kinetics governing resolution of syringomyelia following decompression of the spinal cord remains to be elucidated. In this study, we performed a volumetric analysis of syringomyelia after CM-I decompression and examined the pattern of decay.
Methods

We identified patients with CM-I and syringomyelia who underwent hindbrain decompression. Patients were included for analysis if there were at least 3 postoperative MR images detailing the entire extent of their preoperative syringomyelia, allowing a curve to be plotted. The volume at each time point was calculated by importing DICOM data into OsiriX open-source software (Pixmeo) and identifying the syringomyelia as the region of interest (Fig. 1). Axial coherent oscillatory state acquisition for the manipulation of image contrast (COSMIC) sequences were used for this, allowing for optimal differentiation between the CSF and spinal cord. The 3D volume was rendered and calculated (Fig. 2). We then plotted the syringomyelia volume over time and constructed regression models using the method of least squares. The Akaike information criterion and Bayesian information criterion were used to calculate the relative goodness of fit. The coefficients of determination $R^2$ (unadjusted and adjusted) were calculated to describe the proportion of variability for each individual data set accounted for by the statistical model. An independent biostatistician validated the study methodology and data analysis.

Results

Two patients were identified as meeting inclusion criteria. Plots of the least-squares best fit were identified as $4.01459e^{-0.0180804x}$ for the patient in Case 1 and $13.2556e^{-0.00615859x}$ for the patient in Case 2 (Fig. 3). Decay of the syringomyelia followed an exponential model closely in both patients as evidenced by Akaike information criterion, Bayesian information criterion, $R^2$, and $R^2$-adjusted in Case 1 of 4.77068, 2.92957, 0.989582, and 0.979164, respectively. Case 2 had comparable values of 18.4951, 16.654, 0.948864, and 0.897728, respectively.

Discussion

It is unclear why syringomyelia only develops in a subset of individuals with CM-I. The condition may be related to the degree of compression rather than to the absolute caudal migration of the cerebellar tonsils or to unidentified factors. Other compressive lesions at the cervicomedullary junction can cause syringomyelia. The resolution of syringomyelia after CM-I decompression, however, is well characterized and occurs in the majority of cases. Instances in which syringomyelia fails to resolve postoperatively can be due to arachnoid webbing, extra- or intradural adhesions or scarring, or failure to adequately decompress the craniocervical junction. Interestingly, syringomyelia can also resolve spontaneously on rare occasions. Wetjen and coauthors have estimated the rate of syringomyelia resolution by examining the postoperative images in patients after CM-I decompression. Based on the largest anteroposterior diameter of the syringomyelia, the authors concluded that the median time to greater than 50% narrowing of the syringomyelia was 3.6 months postoperatively, whereas the
Volumetric analysis of syringomyelia

mean time was 6.5 months. To our knowledge, these are the best data available on the rate of syringomyelia resolution after CM-I decompression, although volumetric analysis related to Chiari malformation and interval changes over time go back as far as Nyland and Krogness.13

Most authors describe a syringomyelia by its width or by its cranial to caudal extent. Although measuring its diameter on axial imaging or its longitudinal extent over time allows one to estimate the size of a syringomyelia, it is not a direct measurement of volume. Because of a syrinx being an irregular 3D structure, MR imaging–based volume measurement as suggested in this study may be a more precise measurement than linear dimension estimates. In this study, we plotted the volume of a syringomyelia over time and found that its resolution follows exponential decay. Potential implications of this technique are that, if validated, volumetric resolution of a syrinx may be correlated with the adequacy of hindbrain decompression.

The factors governing the rate of syringomyelia resolution may be complex and depend on the viscosity of the syringomyelia fluid, the shape of the cavity, the elasticity of the spinal cord, and the transmural pressure. Although flow in elastic tubes such as blood vessels has been modeled,19 little is known about the kinetics of syringomyelia. It is likely that the kinetics are more complex than the simple emptying of a container or balloon (Fig. 3). Syringomyelia can take months to empty after surgery, suggesting that complex factors affect this process. Ultimately, the volume of a syringomyelia represents the net difference between factors governing its accumulation and those affecting its drainage, and volumetric measurements over time as in the current study do not allow one to characterize these opposing forces. It appears that emptying of the syringomyelia is not limited by the outflow rostrally through the central canal. If this were the case (simply losing fluid volume through a very narrow central canal), one would expect the bottleneck at the rostral aspect to be the limiting factor and one would see a linear decline in volume over time. In this instance, the exponential decay suggests that the rate of loss is dependent on the residual volume: the greater the residual volume, the greater the rate of egress.

Conclusions

Three-dimensional analysis of syringomyelia resolution following surgery over time enables the kinetics to be estimated and can be described as exponential decay. This analysis should be interpreted as an unvalidated metric that will need to be assessed in a larger group to establish a generalizable mechanism of syrinx resolution. Because syringomyelia is the final common pathway for a number of different pathological processes, it is possible that this exponential decay only applies to syringomyelia related to treatment of CM-I.

Disclosure

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Author contributions to the study and manuscript preparation include the following. Conception and design: Walcott, Coumans. Acquisition of data: Walcott, Coumans. Analysis and interpretation of data: Walcott, Coumans, Butler. Drafting the article: all authors. Critically revising the article: all authors. Reviewed submitted version of manuscript: all authors. Approved the final version of the manuscript on behalf of all authors: Walcott. Statistical analysis: Walcott, Coumans. Study supervision: Coumans.

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