Chiari Malformation: An Overview

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What Is Chiari Malformation?

Chiari Malformation (Arnold-Chiari) - is a serious neurological disorder where the bottom part of the brain, the cerebellum, descends out of the skull and crowds the spinal cord, putting pressure on both the brain and spine causing many symptoms.

- First identified by Hans Chiari in the 1890’s
- Several types, Type I is shown above; Type II is associated with Spina Bifida
- Also known as: Chiari Malformation (CM), Arnold-Chiari Malformation (ACM), tonsillar herniation, tonsillar ectopia, hindbrain herniation

Chiari affects everyone differently!!
Chiari malformation is traditionally defined as the cerebellar tonsils being located 3mm-5mm *or more* below the foramen magnum (opening at the bottom of the skull) as measured on an MRI.

However, research has shown that the size of a herniation is **NOT** strongly related to how severe symptoms are or how people respond to treatment.

Some people have large herniations with no symptoms; while others have small herniations with severe symptoms.
Evolving Definition

Researchers are looking for a new way to measure Chiari severity.

- Focus areas include advanced MRI and engineering techniques to quantify cerebrospinal fluid (CSF) flow, crowding and compliance.
- Some preliminary results are encouraging, but it will take time to develop.

Conquer Chiari is funding research to develop a quantitative, objective test of symptomatic Chiari.
What Causes Chiari?

Cerebrospinal Fluid Issues
Altered CSF dynamics “push” brain out of skull. For example, hydrocephalus, intracranial hypertension.

Small Posterior Fossa
Back of the skull is too small for the normal sized brain. May be most common.

Tethered Cord (very controversial)
Abnormal anchoring of the spinal cord at the base leads to herniation, also causes symptoms of its own.

Other Causes:
- Acquired due to a mass in the brain
- Spina Bifida (Chiari II)
- Genetic conditions such as an Ehlers-Danlos variant
- Incidental; found when an MRI is done for a different reason, such as an accident

It is believed Chiari can be due to several things. However these are only theories and exact mechanisms are not understood. It is not known how common each type may be.
Common Questions

How many people have Chiari?

Chiari is more common than many people think. Conquer Chiari estimates that 1 in 1,000 people have Chiari, which translates to 300,000 people in the US alone. According to the American Association of Neurological Surgeons, there are over 10,000 Chiari surgeries each year.

Who gets Chiari?

It is generally believed that Chiari affects people of all races. There are some indications that it affects more women than men. Symptoms can develop at any age, but people are usually diagnosed as children or as adults in their late 20’s or early 30’s.
Is Chiari genetic?

Research has identified more than 100 families with multiple affected members, implying that at least some cases are genetic in origin. However, it is not known what percent of cases are, and what genes are involved. There is no genetic test for Chiari.

Is Chiari fatal?

In general Chiari 1 is not considered fatal. However, there are case reports of sudden death associated with Chiari 1 due to breathing problems, although this is likely very uncommon. Also, the impact that long-term symptoms may have on life expectancy has not been studied. Finally, severe cases of Chiari II in infants and children can be fatal.
Chiari symptoms can range from mild, to severe, to completely debilitating

- Signature symptom is a severe headache in the back of the head brought on by straining, coughing, etc.
- It is not known why, but symptoms can develop at any age
- Trauma may play a role in sparking symptoms
- Symptoms are due to compression of brain and spinal tissue, disruption of the natural flow cerebrospinal fluid, and increased pressure in the brain
- Some people with malformations are asymptomatic, meaning they have no symptoms
- The cognitive effects of Chiari are being researched, but are not well understood
Syringomyelia

Up to 50% of Chiari patients also develop syringomyelia (SM):

- Neurological condition where a fluid filled cyst, or *syrinx*, forms in the spinal cord
- The syrinx expands the spinal cord, sometimes to more than twice its normal diameter, and stretches the nerve tissue paper thin
- Exactly how Chiari causes a syrinx is not completely understood
  - Likely due to disruption of the natural flow of cerebrospinal fluid (CSF) between the brain and spinal cord
- Can cause permanent nerve damage and paralysis
- Other symptoms include neuropathic pain, loss of sensation, muscle weakness, and bladder/bowel problems
Diagnosis

- There is no single, objective, definitive test for symptomatic Chiari
- Diagnosis is based on a combination of factors and is an informed opinion on whether symptoms are due to herniated cerebellar tonsils
- Chiari is often missed and mis-diagnosed as other diseases

It can take years, and many doctors, for a patient to get an accurate diagnosis
## Treatment

- Only treatment which addresses underlying problem is surgery.
- Treatment decision is based on whether symptoms are clearly due to Chiari and bad enough to warrant surgery.
- No objective test or criteria to say when a patient should have surgery.
- Alternative is to monitor situation and treat individual symptoms (Wait & See).
- Since surgery is the primary treatment option, most patients see a neurosurgeon for treatment opinions.

A 2004 survey of neurosurgeons (Schjiman) found a wide range of thinking on when to operate for Chiari.

<table>
<thead>
<tr>
<th>Case #</th>
<th>Symptoms/Diagnosis</th>
<th>% Would Operate</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>7 yr old with no symptoms 12 mm tonsils- no syrinx</td>
<td>8</td>
</tr>
<tr>
<td>1A</td>
<td>w/ 2mm wide syrinx</td>
<td>28</td>
</tr>
<tr>
<td>1B</td>
<td>w/ 8mm wide syrinx</td>
<td>75</td>
</tr>
<tr>
<td>2</td>
<td>9 yr old with headaches 10 mm tonsils- no syrinx</td>
<td>46</td>
</tr>
<tr>
<td>2A</td>
<td>w/ 2mm wide syrinx</td>
<td>64</td>
</tr>
<tr>
<td>2B</td>
<td>w/ 8mm wide syrinx</td>
<td>90</td>
</tr>
<tr>
<td>3</td>
<td>11 yr old w/progressive scoliosis 12 mm tonsils- no syrinx</td>
<td>58</td>
</tr>
<tr>
<td>3A</td>
<td>w/ small syrinx</td>
<td>85</td>
</tr>
<tr>
<td>3B</td>
<td>w/ 6mm wide syrinx</td>
<td>97</td>
</tr>
</tbody>
</table>

It is common for patients to receive different opinions from different neurosurgeons.
The goal of posterior fossa decompression is to create more room around the malformation and restore the flow of spinal fluid. There are many variations:

**Surgical Steps**

- **Cranietomy** - A piece of the skull is removed
- **Laminectomy** - Part of one or more vertebrae are removed
- **Duraplasty** - The covering of the brain is opened and a patch is sewn in to make it bigger
- **Tissue removal** - Cerebellar tonsils are sometimes cauterized *(controversial)*
- **Plate insertion** - Mesh is inserted where skull was removed

**Surgical Variations/Controversies**

- How much bone is removed
- Whether the dura is opened at all
- What type of material is used for a dural graft (patch)
- Whether any brain tissue is removed
- Whether a mesh is put in to take the place of the removed piece of skull

*Images courtesy of Dr. Ghassan Bejjani*
Surgery - The Experience

Every case is unique, but a typical experience might be:

Surgery: ~ 4 hrs.

ICU: One night

Hospital: 3-4 days

Normal Activity: Variable

Rest at home: 3 wks to 3 mos
Factors that may affect outcomes include: age, syringomyelia, bony abnormalities, scoliosis, duration of symptoms, and concurrent conditions.

Common residual symptoms include pain, muscle weakness, and loss of sensation; patients will adapt their lifestyle accordingly.

Symptoms may fluctuate over time; patients report that trauma can cause symptoms to come back.

Patients in the poor range (10-20%) may end up getting revisions or other surgeries.

Over the very long term (10 years+), symptom recurrence has been reported, but has not been studied extensively.

Published outcomes research has many limitations, such as poorly defined measures and short durations. Patient should discuss their individual case with their surgeon to establish expectations before surgery.
Why Do Surgeries Fail?

*Surgery can fail for a number of reasons*

### Surgery Related
- **Inadequate decompression**
  - not enough bone removed
  - dura was not opened
- **Recurrent obstruction**
  - scarring disrupts CSF flow
  - bone regrowth in children
  - spinal cord retethers
- **Surgical complications**
- **Altered CSF dynamics**
- **Cervical instability**

### Patient Related
- **Concurrent conditions still cause symptoms**
  - pseudotumor cerebri; elevated pressure in the brain
  - basilar invagination; compression of the brainstem from the spine
  - genetic disorders
- **Symptoms not due to Chiari**
  - symptoms were due to something else

It is important to try to understand the reason for failure before undergoing additional treatment, but sometimes a revision surgery will be necessary.
State of Research

**KNOWN**
- Affects >300,000 in the US
- Surgery helps most people
- Can cause syringomyelia
- Affects everyone differently

**QUESTIONS**
- How to identify candidates for surgery?
- Is there an ideal surgical technique?
- What role do genetics play, and what genes?
- Why do people become symptomatic?

**SUSPECTED**
- Several distinct sub-groups/causes
- Children have better outcomes than adults
- CSF dynamics play a role

**CONTROVERSIES**
- The relationship between tethered cord and Chiari
- If children should be restricted from physical activity before and/or after surgery
- If brain tissue should be removed during surgery

Conquer Chiari funds research grants and organizes research conferences to advance our understanding of Chiari and improve experiences and outcomes
Three Keys To Understanding Chiari

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Each person’s Chiari experience is unique

It is impossible for others to completely understand what you are going through; look for validation within yourself.

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Not every question has an answer

Don’t waste time searching for absolutes or a magic bullet.

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There are two options: take control or be controlled

Be proactive, educate yourself, and trust your instincts.
Questions or comments about this presentation can be sent to:
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