

International Beneficent Project
«Colibri»
for the disabled children diagnosed with
The Pseudobulbar Palsy



Why the Project is titled «Colibri»?

The unusually long beak of the colibri reminds an enteral feeding tube through which children diagnosed with Pseudobulbar Palsy eat



What does the diagnosis mean?

Children Pseudo bulbar Palsy -

is a neurologic disorder. One of its reason is the birth injury. Particularly nasty form of the Pseudo bulbar Palsy, along with other features, characterized by the inability to eat by mouth. Chewing and swallowing is the lethal risk for such children because food enters the respiratory tract.

Throughout life The children don't know what a food taste is.

It's unbelievable and unimaginable!

Assumed that

birds fly only beak forward



**Children Pseudobulbar Palsy
is the inextirpable disease**



However, it is known that

The tiny Calibri can fly by tail forward



Many diseases, previously referred to the incurable, today are successfully treated



**"Seek and ye shall find;
knock, and it shall be
opened to you"
(St. Matthew 7,7)**

Why The Project is important?



Medicine has learned methods to save the lives of children with serious birth defects consequence of which is The Pseudo bulbar Palsy.

Subsequently, children with this diagnosis can not receive the full treatment and rehabilitation due to lack of systematic experience in this area.

The Pseudo bulbar Palsy
is referred to the inextirpable disease.

But the current level of medicine allows you to review this statement!

The aim of The Project «Colibri»

Attract the attention of the public, the medical and scientific communities to the problems of treatment, rehabilitation and the quality of the daily life of children diagnosed with The Pseudobulbar Palsy.

The objectives of The Project «Colibri»



- 1.To create a website, to answer questions of treatment, rehabilitation and the daily life of children diagnosed with The Pseudobulbar Palsy.
2. To provide practical assistance to families with children suffering from diagnosed The Pseudobulbar Palsy.

We are the little heros. We had endured the long months of reanimation when we tried to learn how to breathe. We endured the excruciating pain. Thank God! Thank doctors!

Thanks to everyone who was with us in the most difficult moments!

It's time to make a new step!

Now it's dangerous to us to eat by mouth, but we dream to eat like normal people.

We want to feel the taste of food.

We ask you to help us!

The Boy Tuktabaev Rustam, 9 years
Tyumen Region



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Who saves one life, saves the whole world!



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