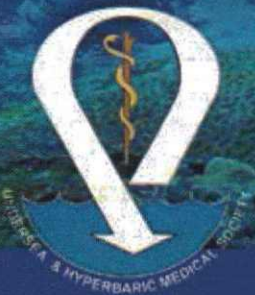
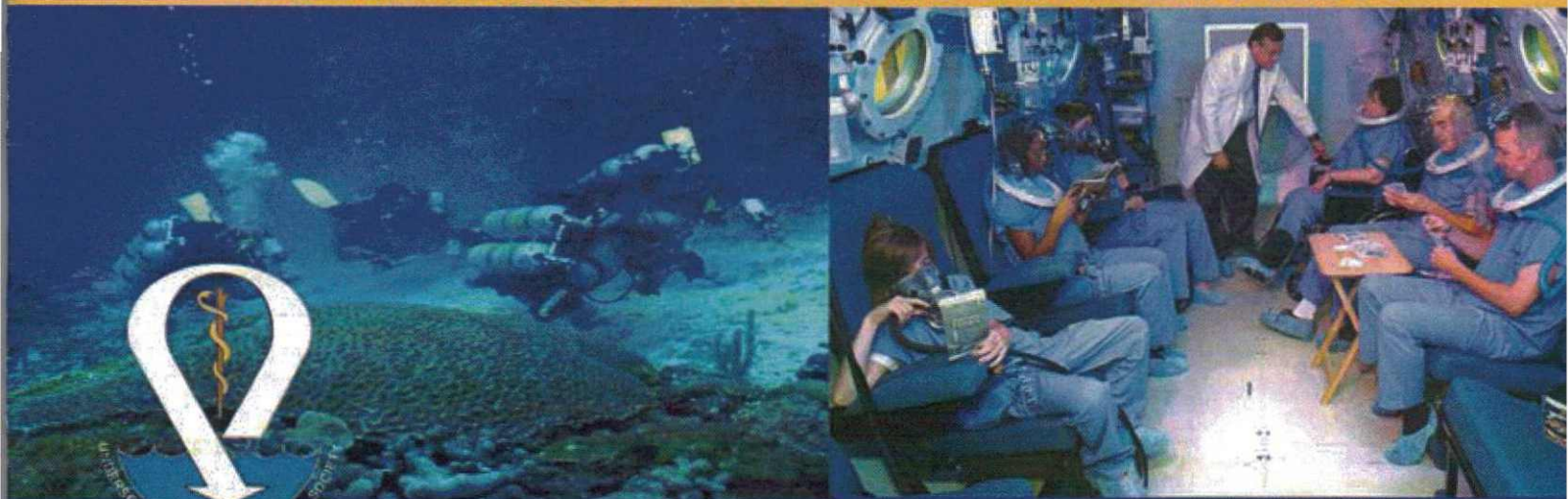


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SYNOPSIS

In April 2014, a controlled longitudinal study to assess the effects of intensive rehabilitation combined with hyperbaric oxygen therapy (HBOT) on gross motor function in 150 children with cerebral palsy (CP) was published in the Undersea Hyperbaric Medical Society (UHMS). This is the first study that compared the effects of different hyperbaric dosages combined with rehabilitation in children with CP, to a control group who received only rehabilitation. In addition, this study is the first empirical study recognized by the UHMS that demonstrated the efficacy for mild-pressure HBOT. Furthermore, this study has shown that a therapeutic effect can be achieved without administering 100% oxygen; for which compressed room air provided nearly identical results.

The eight-month study commenced after assigning the children into four groups. The control group consisted of children who did not undergo hyperbaric treatments and participated in intensive rehabilitation only. The three other groups were assigned to hyperbaric treatment, in addition to receiving intensive rehabilitation. The rehabilitation program was applied for two hours per day, six days a week over six months and consisted of a half-hour of individual therapies including physical therapy, occupational therapy, speech therapy and special education. All three HBOT groups received 40 one hour sessions, six days a week. Every child received the same intensive rehabilitation care by the same therapist team, at the same center, using the same protocol and the same duration of follow-up. The first HBOT group was treated at 1.3 atmospheres absolute (ATA) with compressed air (21% oxygen). No supplemental oxygen was given to this group. The second HBOT group was treated at 1.5 ATA with 100% oxygen delivered through a hood system. Finally, the third HBOT group was treated at 1.75 ATA, also with 100% oxygen delivered through a hood system.

As expected with intensive therapies, all four groups improved. However, the findings demonstrated that the three groups treated with different dosages of HBOT improved significantly more than the control group. The HBOT groups' gross and fine motor (GMFM) function variations were on average 3 times higher than the control.

For the three HBOT groups, there was no significant statistical difference between the observed improvements. The study found that administering mild-pressure HBOT at 1.3 ATA has the potential to be equally effective when compared to higher pressures for children with CP. Furthermore, the U.S. Agency for Healthcare Research and Quality (AHRQ) analyzed the results of this study and arrived at the same conclusions. The AHRQ report said, "The possibility that pressurized room air had a beneficial effect on motor function should be considered the leading explanation." In this context, the increase of upwards to 50% plasma oxygenation achieved by compressed air at 1.3 ATA was of significance. This study provides further credibility to the substantial effects of HBOT at mild-pressure.

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Intensive rehabilitation combined with HBO₂ therapy in children with cerebral palsy: A controlled longitudinal study

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ABSTRACT

Objective: The present study aimed to assess the effect of intensive rehabilitation combined with hyperbaric oxygen (HBO₂) therapy on gross motor function in children with cerebral palsy (CP).

Methods: We carried out an open, observational, platform-independent study in 150 children with cerebral palsy with follow-up over eight months to compare the effects of standard intensive rehabilitation only (control group $n = 20$) to standard intensive rehabilitation combined with one of three different hyperbaric treatments. The three hyperbaric treatments used were:

- air (FiO₂ = 21%) pressurized to 1.3 atmospheres absolute/atm abs ($n = 40$);
- 100% oxygen pressurized at 1.5 atm abs ($n = 32$); and
- 100% oxygen, pressurized at 1.75 atm abs ($n = 58$).

Each subject assigned to a hyperbaric arm was treated one hour per day, six days per week during seven weeks (40 sessions). Gross motor function measure (GMFM) was evaluated before the treatments and at two, four, six and eight months after beginning the treatments.

Results: All four groups showed improvements over the course of the treatments in the follow-up evaluations ($p < 0.001$). However, GMFM improvement in the three hyperbaric groups was significantly superior to the GMFM improvement in the control group ($p < 0.001$). There was no significant difference between the three hyperbaric groups.

Conclusion: The eight-month-long benefits we have observed with combined treatments vs. rehabilitation can only have been due to a beneficial effect of hyperbaric treatment.

INTRODUCTION

Cerebral palsy (CP) is due to a lesion of the developing brain, characterized by inadequate muscle tone and control, often associated with other types of neurodevelopmental delay involving cognitive, communication and psychosocial skills. Treatments are mainly focused on exploiting residual cerebral function, and intensive rehabilitation is recognized to have demonstrated its efficacy in achieving better function and autonomy, thus creating a better quality of life [1].

The leading causes for cerebral palsy stem from a critical reduction of oxygen (O₂) delivery to a part of the developing brain in the perinatal period [2]. The site of the brain lesion can be localized with cerebral blood

flow measurements using brain single-photon emission computerized tomography (SPECT) [3,4] because impaired brain cell nutrition and oxygen delivery are related to inadequate blood flow. While hypoxia may cause neuronal death, there is a well-known phenomenon called the “ischemic penumbra,” which defines a volume of tissue surrounding a zone of infarction where cells receive enough oxygen to survive in an “idling state,” but not enough to function normally [5]. It has been suggested that these neurons might be viable much longer than previously believed [6,7,8], and this is where regenerative medicine is trying to play a role. Hyperbaric oxygen (HBO₂) treatment has shown reproducible benefits for more than two decades in hundreds of

children with CP around the world [9]. Using high-quality SPECT imagery, several studies of children with CP and of adults after a stroke have shown that HBO₂ therapy may regenerate or revive cells in the ischemic penumbra in the brain [7,10,11]. This increased vascular activity would allow the reactivation of “idling” neurons [6,10,11, 12], as HBO₂ therapy is known to increase neovascularization in wound healing. The higher tissue oxygen levels provided by HBO₂ therapy might also favor better metabolism and function of unaffected cells [13,14].

To date, despite several reports of benefit, the use of HBO₂ therapy for CP has met opposition, which has even polarized the field of clinical HBO₂ therapy [15-18]. The first pilot study [19] reported the positive effects of HBO₂ therapy on 25 carefully selected children with the form of CP known as spastic diplegia. The improvements were measured both on gross and fine motor function. Based on the results of this pilot study, a double-blind randomized multicenter trial ($n = 111$) of HBO₂ therapy for children with CP was conducted by Collet *et al.* [20]. This study included only two groups of children: one treated at 1.75 atmospheres absolute (atm abs) with 100% O₂, while the other breathed air at 1.3 atm abs. Some involved in the statistical analysis of the results regarded the use of compressed air at 1.3 atm abs to be an inactive placebo, although this was opposed by the clinicians.

The controversy required the appointment of an independent adjudicator by the *Lancet*, who agreed that such a change in pressure and increase in the level of oxygen could not be referred to as a “sham” treatment. In fact, exposure to 1.3 atm abs increases the arterial plasma oxygen concentration (PaO₂) by nearly 50% [21]. It was little recognized at the time that blood flow in the physiological range of oxygen concentrations is controlled by the interaction between nitric oxide and hemoglobin [22]. Changes in oxygen levels also regulate genes involved in angiogenesis and neutrophil activity in inflammation [23]. As the best dosage of oxygen for the treatment of children with CP is not known, a sham control group should have been included to ensure an adequate experimental design. The controversy was highlighted by an editorial comment entitled “Hype or hope” published in the same issue of the *Lancet* journal [24].

After the courses of treatment, the improvements on gross motor function were impressive and equivalent in both groups. Improvements in language and neuropsychological functions were also recorded in both treatment groups. There are two ways of interpreting the

results: either the two treatments were equally effective, or the improvements were all caused by a “participation effect.” Based on the major improvements reported, the latter interpretation is inappropriate [25] but has, unfortunately, been promoted as evidence that hyperbaric treatment is ineffective in CP children [26] restricting further research on the subject. The aim of this present study is to answer the questions raised by the study by Collet *et al.* [20] by assessing the effect of different dosages of hyperbaric treatment combined with intensive rehabilitation on motor function in children with CP.

METHODS

Participants

A total of 150 children with CP were selected for the study among those attending rehabilitation at the Foundation for Spastic and Mentally Handicapped Persons-UDAAN (FSMHP-UDAAN) center in Delhi, India. All participants had to meet the following inclusion criteria: children up to teen age of either sex with all types of CP, any cognitive and motor development level.

Children were excluded if there were other developmental or genetic disorders, uncontrolled epilepsy or asthma, as well as ear, nose or throat disorders. Forty percent of all of our participants had minor to moderate epilepsy due to their injured brain. Half of them were significant enough to be on antiepileptic medication. It was the parents’ decision to include their children in the HBO₂ therapy groups. Participants who were not assigned to HBO₂ therapy groups were assigned to the control group. All participants were engaged in the same intensive rehabilitation program at FSMHP-UDAAN. Only the children who did not default on at least six months of standard therapies were assessed. Quality, magnitude and type of care were uniform across all four groups. Participants’ characteristics are described in Table 1. The study was approved by the ethics committee of Apollo Hospital, Delhi, and the parents’ informed voluntary written consent was required after medical clearance.

Treatments

The study covers a 10-year span of treatments during which the three different dosages of hyperbaric oxygen were used. The different dosages were not implemented at the same time, and the children were offered the HBO₂ therapy available at the time of their inclusion in the protocol, which means that no selection bias occurred in the choice of dosage.

Table 1: Participants' characteristics

Groups	Diagnostics	Gender (M/F)	Age (yrs) Mean (range)	GMFM
				baseline score Mean (SD)
Control (n=20)	Athetoid CP, n=2 Hemiplegic CP, n=2 Diplegic CP, n=4 Quadriplegic CP, n=12	13/7	3.5 (1 to 17)	29.6 (13.0)
1.3 atm abs (n=40)	Athetoid CP, n=3 Hemiplegic CP, n=0 Diplegic CP, n=16 Quadriplegic CP, n=12	29/11	4.9 (1 to 11)	29.6 (14.8)
1.5 atm abs (n=32)	Athetoid CP, n=3 Hemiplegic CP, n=1 Diplegic CP, n=15 Quadriplegic CP, n=13	23/9	4.3 (1 to 12)	34.3 (15.6)
1.75 atm abs (n=58)	Athetoid CP, n=6 Hemiplegic CP, n=2 Diplegic CP, n=19 Quadriplegic CP, n=31	40/18	4.3 (1 to 13)	32.5 (11.8)

atm abs = atmosphere absolute; CP = cerebral palsy; F = female;
GMFM = gross motor function measurement; M = male.

Every child received the same intensive rehabilitation care by the same therapist team, at the same center, using the same protocol, and the same duration of follow-up. The rehabilitation program was applied for two hours/day, six days/week over six months, and consisted of a half-hour of individual therapies of physical therapy, occupational therapy, speech therapy and special education.

For hyperbaric therapy, the children were assigned to four groups:

- A- No hyperbaric treatments, rehabilitation only (control group), n=20;
- B- 40 sessions, one hour/day, six days/week at 1.3 atm abs air, 21% O₂ (room air), n=40;
- C- 40 sessions, one hour/day, six days/week at 1.5 atm abs HBO₂, 100% O₂, n=32;
- D- 40 sessions, one hour/day, six days/week at 1.75 atm abs HBO₂, 100% O₂, n=58.

All hyperbaric treatments were given six days/week during seven weeks. In all treatment sessions, the total amount of time spent in the hyperbaric chambers was 90 minutes, as 15 minutes for either compression and decompression was taken. HBO₂ using 100% oxygen was delivered through a hood inside a multiplace hyperbaric

chamber at a local tertiary care hospital, using pressures of 1.75 or 1.5 atm abs. Hyperbaric air treatment at 1.3 atm abs using room air at 21% oxygen was carried out using a soft chamber. We carried out initial and periodic assessment of lung and ENT passages and temporarily stopped hyperbaric therapy whenever there was any air passage obstruction or inflammation. Children with a previous history of epilepsy were referred to a pediatric neurologist, and the anti-epileptic dosages were increased marginally during the hyperbaric treatments period.

Evaluation procedures

In all children, gross motor function was systematically evaluated before the treatments and at four and six months after the beginning of the treatments by the same therapists, who were accustomed to undertaking the evaluations. To have more data, and when possible, we were often able to evaluate the children at two and eight months after the beginning of treatments. The gross motor function measure (GMFM66) [27] was applied to every child. It is a criterion-based observational measure (66 items) that assesses motor function in five dimensions: A-lying and rolling, B-sitting, C-crawling and kneeling, D-standing and E-walking, running and jumping.

Table 2: GMFM observed mean before and after HBO₂ therapy

	<i>GMFM observed mean (SD)</i>				
	Before HBO ₂	2 months after beginning HBO ₂	4 months after beginning HBO ₂	6 months after beginning HBO ₂	8 months after beginning HBO ₂
Control	29.6 (13.0)		31.0 (12.8)	32.4 (12.8)	
1.3 atm abs 21% O ₂	29.6 (14.8)	33.4 (13.1)	36.2 (13.6)	38.6 (14.3)	40.8 (14.2)
1.5 atm abs 100% O ₂	34.3 (15.6)		39.3 (15.4)	42.5 (15.3)	46.4 (17.0)
1.7 atm abs 100% O ₂	32.5 (11.8)		37.2 (10.8)	42.1 (10.4)	46.7 (9.7)

atm abs = atmosphere absolute; GMFM = gross motor function measurement

Each item is scored on a four-point scale, and the test gives numeric results for each dimensions as well as a total score. The score is reported as a percentage of the maximum score (100%) generally obtained in a normal 5-year-old child.

Data analysis

Linear mixed models were used to analyze the GMFM data. Such models permit the data to exhibit correlations and non-constant variances. These models, therefore, provide the flexibility of modeling not only the means of the data but also their variances and co-variances. Treatments were considered as fixed factors, and month and age were considered as co-variables. Month was time-dependent, while age was time-independent. Random components were introduced to depict individual trajectories over months with separate intercepts and slopes. A maximum likelihood approach was used to estimate the coefficients, and an unstructured random effect covariance matrix was utilized. Linearity for month and interactions (treatment \times month) were tested. Information criteria (such as the Akaike criterion and the $-2\ln$ (likelihood)) and residual values were used to verify the quality of adjustment. Pearson product-moment correlation coefficient (r) was calculated to quantify the interrelationship among the GMFM variation and GMFM level before HBO₂ therapy.

RESULTS

As expected, groups were similar on the GMFM level at baseline ($p = 0.429$) and each group, including the control group, showed improvement in the GMFM scores over the course of the treatments ($p < 0.001$). As depicted in Table 3, there were statistically significant interactions between group and month ($p < 0.001$) and a statistically significant age effect ($p = 0.003$). To better

understand these results, fixed-effect linear models are presented in Table 4 for each group. We observe that the GMFM score increases by 0.46 unit per month in the control group as compared to values ranging from 1.36 to 1.50 unit per month in the experimental groups; and these slopes are significantly different from the control group slope ($p < 0.001$). These results are visualized in Figure 1. GMFM variation, which is the average monthly improvement in the GMFM results over the course of the follow-up, was correlated with GMFM level before HBO₂ therapy ($r = -0.33, p < 0.001$).

DISCUSSION

This is the first study that has compared the effects of different hyperbaric dosages combined with rehabilitation in children with CP to a control group receiving only rehabilitation. As expected with intensive therapies, all four groups improved substantially. However, our findings demonstrate that the three groups treated with different dosages of HBO₂ improved much more than the control group, as their GMFM variations were on average three times higher.

In the present study, the three treatments were equally effective in producing gross motor improvement. This reproduces the impressive results obtained in the two groups (1.5 atm abs HBO₂, 100% O₂ and 1.3 atm abs air) in the study of hyperbaric treatment for CP children by Collet *et al.* [20]. Mychaskiw has pointed out in a UHM editorial that children treated with compressed air at 1.3 atm abs cannot be regarded as a control group [28]. It is obvious that giving more oxygen for neurologic conditions is not an all-or-none phenomenon. We find it disconcerting that such a flawed study has been used to claim a lack of efficacy of hyperbaric treatment in cerebral palsy when Collet *et al.* [20] actually stated: "The improvements in GMFM scores in both groups are

Table 3: Fixed effects estimation for GMFM

Variable	Coefficient (B)	SE(B)	T	p-value
Constant	24.65	3.31	7.45	0.000
1.3 atm abs	-1.91	3.65	-0.52	0.602
1.5 atm abs	2.91	3.73	0.78	0.437
1.75 atm abs	1.42	3.39	0.42	0.675
Month	0.46	0.18	2.52	0.013
LnAge	4.96	1.66	2.99	0.003
1.3 atm abs* month	0.90	0.22	4.14	0.000
1.5 atm abs* month	0.94	0.23	4.16	0.000
1.75 atm abs* month	1.04	0.210	4.95	0.000

atm abs = atmospheres absolute; GMFM = gross motor function measurement

Table 4: Predicted GMFM from fixed effects models in each group

Group	Model
Control group	GMFM = 24.65 + 0.46 Month + 4.96 LnAge
1.3 atm abs group	GMFM = 22.75 + 1.36 Month + 4.96 LnAge
1.5 atm abs group	GMFM = 27.56 + 1.40 Month + 4.96 LnAge
1.75 atm abs group	GMFM = 26.07 + 1.50 Month + 4.96 LnAge

atm abs = atmospheres absolute; GMFM = gross motor function measurement

clinically important... The improvement seen in all other outcomes is also striking." Moreover, the U.S. Agency for Healthcare Research and Quality (AHRQ) analyzed the results of the study and arrived at the same conclusions [25]. The AHRQ report mentioned that "The possibility that pressurized room air had a beneficial effect on motor function should be considered the leading explanation."

However, our study has, like that of Collet *et al.* [20], clearly demonstrated the benefit of treatment with compressed-air at 1.3 atm abs, because we included a control group; thus the effect of hyperbaric conditions cannot be attributed to a participation or placebo effect. In fact, the placebo effect is a temporary phenomenon that lasts for a few weeks [29] and not for the eight months we have found benefit in our follow-up. Human physiology works within a narrow band for optimal activity. In this context, the increase of almost 50% in plasma oxygenation achieved by compressed air at 1.3 atm abs was of significance.

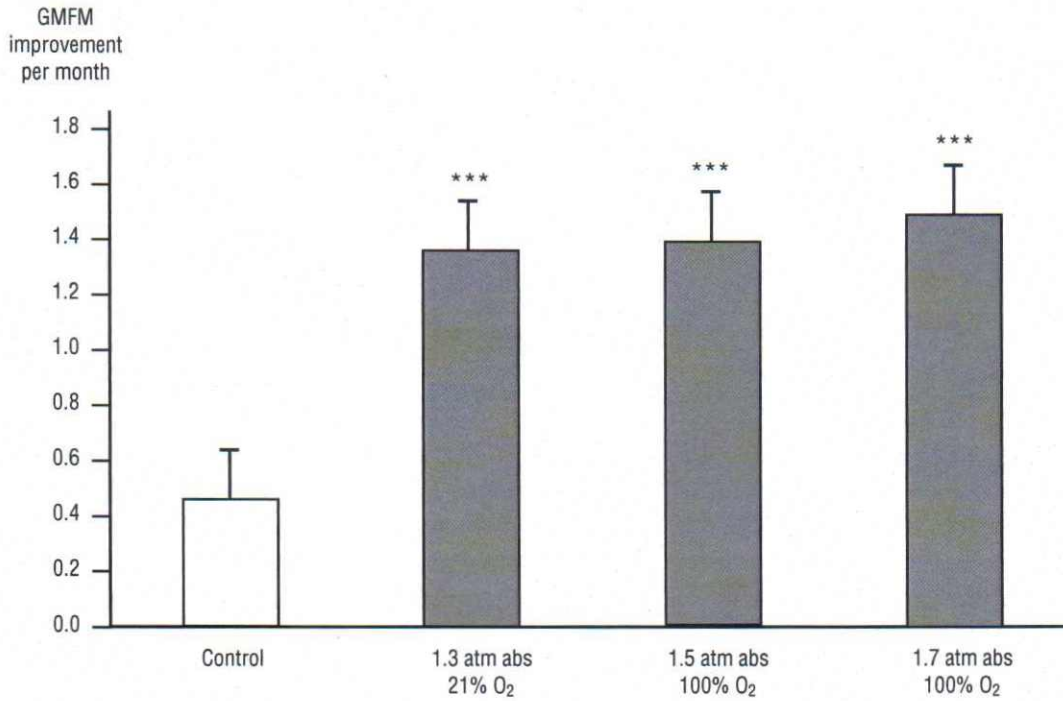
A study on patients with advanced lung disease has been undertaken in Jerusalem. While maintained on supplemental oxygen, they were taken down to the Dead Sea, where they breathed only ambient air. A statistically significant increase in walking distance was recorded, which persisted for a month after returning to Jerusalem. The increase in pressure achieved by descending to the Dead Sea was just 0.06 atm abs [30]. Compressed air at a pressure 0.3 atm abs over ambient cannot be considered a placebo; and a recent paper discussed the osmotic effects of a sudden increase in pressure [31]. In addition, most of the children included in our series were barely in a position to have the mental maturity to understand what was being done for them.

The results of the present study strongly support the fact that HBO₂ therapy, even in small dosage, can improve motor function and increase the effects of standard rehabilitation. The amount and quality of changes observed in our study are also in accordance with the results obtained in other studies on HBO₂ therapy in CP [10,19,20]. The authors are aware that Lacey *et al.* [32] have recently conducted a randomized control study in which they compared two different hyperbaric treatments, one of which (14% O₂ at 1.5 atm abs) has never been used on CP children before, and was considered by these authors as a control group. These authors present their study as a definitive answer to hyperbaric therapy inefficacy in children with CP even if major concerns can be addressed and explain the discrepancy with the present study.

First, despite the fact that in the control group, the condition simulated 21% oxygen at room air, this treatment must not be considered as a placebo treatment because no one knows exactly the potential physiologic effects of this hyperbaric treatment. Secondly, the change in GMFM in the HBO₂ group was 1.5 in two months, which is more than most changes measured with recognized treatments in CP [9]. Thirdly, Lacey *et al.* included only 20 participants per group and stopped the study prematurely, which avoided possibility of the results reaching a threshold for significance. These concerns have been addressed in a letter to the *Annals of Neurology* [33].

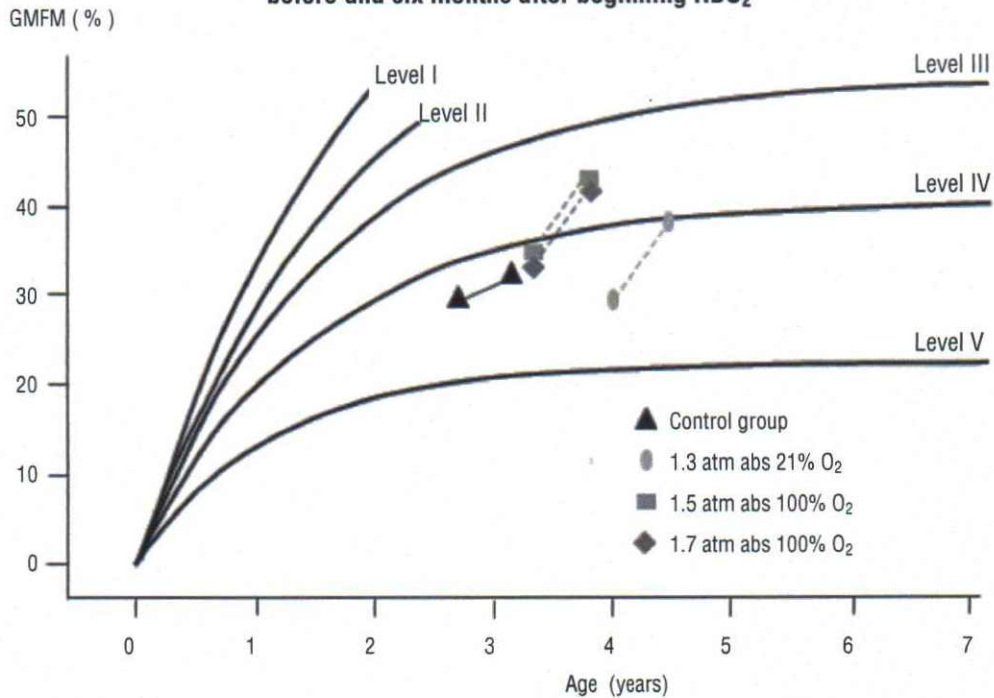
The Gross Motor Function Classification Scale (GMFCS) classifies CP disabilities into five levels based on the GMFM measurement at a given age. The natural gross motor progression of children with CP usually

Figure 1: Rate of gross motor function measurement improvement



*** = significantly different from the control group, $p < 0.001$; atm abs = atmospheres absolute

Figure 2: Gross motor function classification scale values before and six months after beginning HBO₂



atm abs = atmosphere absolute; GMFM = gross motor function measurement

follows a curve similar to a logarithmic curve [27]. The children with the highest level of abilities are classified in Level 1, while Level 5 regroups the children with the most severe form of motor disability (Figure 2). The progression of children with CP should naturally follow the curves corresponding to their level of disabilities [27]. Figure 2 shows that the mean initial GMFM values of the four groups would classify them between Level 4 and Level 5 of the GMFCS. By end of six months of therapy, all three hyperbaric groups had improved to Level 4, whereas the control group did not change its disability level.

There are risks associated with the high oxygen pressures used in diving, but they are not relevant to the much lower pressures used in this study. The rate of change of pressure was slow, as the pressurization took 15 minutes, and only three children were excluded because of ear pain on compression. None of the participants needed ear canal grommet use. There were no other side effects.

Our study shows that HBO₂ therapy, when combined with rehabilitation, has many more positive effects than rehabilitation alone. As seen on SPECT imaging, hyperbaric treatment appears to reactivate certain damaged areas of the brain. It is, however, obvious that the recovering brain must be trained to work to its full potential to gain the best results. This highlights the importance of rehabilitation after or during HBO₂ therapy. Further research is needed to explore the cerebral plasticity processes that follow hyperbaric treatment. Improvement in function, comfort and the independence of children with disabling neurological conditions could lead to better health and quality of life as well as important cost savings in the long term.

LIMITATIONS

There were several limitations inherent to this study. First, participant repartition between groups was not randomized. It was the parents' decision to include their children in HBO₂ therapy groups, and participants who were not assigned to HBO₂ groups were automatically assigned to the control group. The different dosages of HBO₂ were not implemented at the same time over a 10-year period, which means that no selection bias occurred in the treatment or dosage choice.

Secondly, the evaluations were not blinded. We certainly recognize that it was not ideal, but it was difficult for us, in a longitudinal study conducted in a relatively small center and involving human interaction and evaluation by the same therapists, for blinded evaluations to have been undertaken.

CONCLUSION

A longitudinal study in children with cerebral palsy has been conducted. The study compared three different dosages of hyperbaric oxygen, combined with intensive rehabilitation with a control group receiving only rehabilitation. The rate of improvement in GMFM score was significantly superior in the three hyperbaric groups compared to the control group. There was no difference between the three HBO₂ therapy groups. The amount of changes are similar to the results obtained in the multiple studies on HBO₂ therapy in CP that have been published and are more important than the improvements measured with standard recognized therapies alone in CP. The very important difference observed in treated vs. controlled children can only be a genuine beneficial effect of HBO₂ therapy. Based on the results of this and other studies of HBO₂ therapy in CP children, HBO₂ combined with rehabilitation should be recommended for children with CP.

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The authors report that no conflict of interest exists with this submission. ■

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