



One life. One opportunity.

Congratulations on your pregnancy!

Your little miracle will soon make your world more beautiful and happier. As expecting parents, you must be busy preparing for your little bundle, ensuring that you give your baby nothing but the best.

Health is wealth and you now hold the key to providing a lifetime of good health for your baby by preserving the babies stem cells at the time of birth. Stem cell collection and preservation is as easy as flipping the pages of a book. As you learn more about the importance of stem cells and their preservation through this journey, we hope that you **make a responsible choice today and secure your child's and family's health for a life time.**

All about stem cells

The umbilical cord is the magical bond between the mother and the baby, whilst providing nourishment to the baby in the womb also plays a key role in protecting your family for a lifetime.

The umbilical cord is a rich source of Haematopoietic (blood forming) Stem Cells (HSC's) and Mesenchymal (tissue and organ forming) Stem Cells (MSC's). Stem cells are master cells that possess the ability to specialise and are thus used as the basic building material in the body's constant renewal process.

The HSC's are being successfully used to cure diseases like Leukaemia, Thalassemia, Sickle Cells Anaemia and more such 80+ life threatening disorders. More than 50,000 stem cell transplants have already been performed worldwide and ~ 1540 in India*. MSC's have potential to facilitate repair, regenerate and restore normal tissue function. More than 362 clinical trials using MSC's are currently in progress including treatment for Cerebral Palsy, Autism and Type 1 Diabetes**.

The umbilical cord also has ability in treating chronic diseases like hearing loss and AIDS with multiple trials in process.

The stem cells found in the cord blood and tissue are genetically unique to your baby and family. As a parent you should consider opting for cord blood stem cell preservation to protect your family from 80+ serious diseases.

*http://www.who.int/transplantation/hscx/en/

**http://bit.ly/2uuvZz9



Since 1988, doctors have been using cord blood in lifesaving treatments. Cord blood stem cells are now being successfully used to treat conditions like Leukaemia and Thalassemia in India. The future of stem cell technology is promising and will save many lives.

Dr. Vijay Ramanan, MD, DM

Leading Clinical Haematologist specialising in Stem Cell Transplantation (Pune, India)



100+ treatable diseases

Cord blood stem cells can treat 90+ life threatening diseases:

Leukemia

- Acute Lymphoblastic Leukemia (AII)
- Acute Myelogenous Leukemia
- Acute Biphenotypic Leukemia
- Acute Undifferentiated
- Chronic Lymphocytic Leukemia (CLL)
- Chronic Myelogenous Leukemia
- Juvenile Chronic Myelogenous Leukemia (JCML)
- Juvenile Myelomonocytic Leukemia (JMML)

Myelodysplastic Syndromes

- Refractory Anemia
- Refractory Anemia with Ringed Sideroblasts (Sideroblastic
- Refractory Anemia with Excess Blasts

Lymphoma

- Hodgkin's Lymphoma
- Refractory Anemia with Excess Blasts in Transformation
- Chronic Myelomonocytic Leukemia (CMML)
- Non-Hodgkin's Lymphoma (Burkitt's Lymphoma)

Other Disorders of Blood Cell Proliferation

- Aplastic Anemia
- Fanconi Anemia
- Congenital Dyserythropoietic Anemia
- Paroxysmal Nocturnal Hemoglobinuria
- Sickle Cell Disease
- Beta Thalassemia Major
- Diamond-Blackfan Anemia
- Pure Red Cell Aplasia
- Amegakaryocytosis / Congenital Thrombocytopenia
- Glanzmann Thrombasthenia

Other blood complications

- Amyloidosis
- Dyskeratosis Congenita
- Myelodysplastic Syndrome

Inherited Immune System Disorders: Severe Combined Immunodeficiency

- SCID with Adenosine Deaminase Deficiency (ADA-
- SCID which is X-linked
- SCID with absence of T & B Cells
- SCID with absence of T Cells. Normal R Cells

Omenn Syndrome

Inherited Immune System Disorders

- Infantile Genetic Agranulocytosis (Kostmann Syndrome)
- Myelokathexis
- Congenital Neutropenia
- Ataxia-Telangiectasia
- Bare Lymphocyte Syndrome
- Common Variable Immunodeficiency
- DiGeorge Syndrome
- Hemophagocytic Lymphohistiocytosis
- Leukocyte Adhesion DeficiencyLymphoproliferative Disorders

- Lymphoproliferative Disorder, X-linked
- Wiskott-Aldrich Syndrome
- Cartilage-Hair Hypoplasia
- Erythropoietic Porphyria
- Hermansky-Pudlak Syndrome
- Pearson's Syndrome
- Shwachman-Diamond Syndrome
- Systemic Mastocytosis
- **Fucosidosis**
- Thymic Dysplasia
- X-linked Agammaglobulinemia

Other Immune Disorders

- **Evans Syndrome**
- Hemophagocytosis Langerhans' Cell Histiocytosis (Histiocytosis X)

HIV - AIDS

Myeloproliferative Disorders

- Acute Myelofibrosis
- Agnogenic Myeloid Metaplasia (Myelofibrosis)
- Polycythemia Vera
- Essential Thrombocythemia

Phagocyte Disorders

- Chediak-Higashi Syndrome
- Chronic Granulomatous Disease
- Neutrophil Actin Deficiency
- Reticular Dysgenesis

Bone Marrow Cancers

- Multiple Myeloma
- Plasma Cell Leukemia
- Waldenstrom's Macroglobulinemia

Inherited Metabolic Disorders

Mucopolysaccharidosis (MPS) Storage Diseases

- Hurler Syndrome (MPS-IH)
- Scheie Syndrome (MPS-IS)
- Hunter Syndrome (MPS-II)
- Sanfilippo Syndrome (MPS-III)
- Morquio Syndrome (MPS-IV)

Metachromatic Leukodystrophy

Pelizaeus-Merzbacher Disease

- Maroteaux-Lamy Syndrome (MPS-VI)
- Sly Syndrome (MPS-VII)
- Mucolipidosis II

Leukodystrophy Disorders

- Adrenoleukodystrophy (ALD)
- Krabbe Disease

Lysosomal Storage Diseases

- Niemann-Pick Disease
- Sandhoff Disease

Wolman Disease Mannosidosis

- Other Inherited Metabolic Disorders
- Lesch-Nyhan Syndrome Osteopetrosis
- Congenital Erythropoietic
- Porphyria (Gunther Disease) Gaucher Disease

Cancer

Solid tumors not originating in the blood or immune system

- Neuroblastoma
- Medulloblastoma
- Retinoblastoma
- **Others**
- Hearing loss
- Hypoplastic left heart syndrome
- Chronic Active Epstein Barr





Giving New Hope



Cerebral Palsy

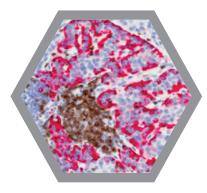
Cerebral Palsy is the result of a brain injury or a brain malformation in the womb. It affects around 10,000 infants annually worldwide. CP affects muscles and a person's ability to control them.

Umbilical cord stem cells have shown potential in the treatment of CP. In a study, two toddlers suffering from CP underwent autologous umbilical cord blood (UCB) transfusion. Gross Motor Function Classification System (GMFCS) improvements were seen in both without any side effects thereby encouraging functional outcome improvements in children with CP.

Type 1 Diabetes

Type 1 Diabetes, also known as juvenile diabetes, is usually diagnosed in children and young adults. In this diabetes, the body does not produce insulin.

In a study, twenty adult patients with newly diagnosed type 1 diabetes underwent MSC treatment and showed tremendous changes without any side effects. Autologous MSC transplantation in new-onset type 1 diabetes constituted a safe and promising strategy to intervene in disease progression and preserve β -cell function.



*https://www.ncbi.nlm.nih.gov/pubmed/25204974?dopt=Abstract



Congenital Heart Disease

It is also known as congenital heart anomaly which is a problem in the structure of the heart that is present at birth. Every year in India, around 78,000 infants die of Congenital Heart Disease*.

In a study conducted by S Tarui, 14 patients suffering from hypoplastic left heart syndrome (HLHS) were injected with c**ardiac progenitor cells** and no complications, including tumor formation, were reported within 36 months. This therapeutic strategy may enhance somatic growth and reduce incidence of heart failure.

Total Hearing Loss

Hearing loss is a common problem that often develops with age or is caused by repeated exposure to loud noises. About 63 million people in India suffer from acquired Hearing Loss.

A success story was observed where the use of umbilical cord blood stem cells has treated a child with Hearing Loss*.

Stephanie Connor acquired a virus infection during her pregnancy which had put her daughter at significant risk for brain damage and lifelong Hearing Loss at age 1.

In January 2012, Madeleine, 2, became the first child to undergo an experimental Hearing Loss treatment through stem cells from her own banked cord blood into her damaged inner ear. Madeleine has since then seen dramatic improvement in her ability to hear.

*Reference: http://abcnews.go.com/Health/Wellness/cord-blood-stem-cells-restore-hearing-toddler/story? id=16750718



The above case studies offer a compelling reason why parents should consider preserving their child's cord blood, since it may offer a treatment option in the future



^{*}https://www.ncbi.nlm.nih.gov/pubmed/21335665

 $[\]verb|^*https://www.ncbi.nlm.nih.gov/pubmed/26232942?dopt=Abstract|\\$



Hospitals that have reported heamatopoitic stem cell transplants

- Christian Medical College (CMC), Vellore
- Tata Medical Center, Kolkata
- Apollo Specialty Hospital, Chennai
- Narayana Health, Bangalore
- Medanta-The Medicity, New Delhi
- BL Kapoor, New Delhi
- Deenanath Mangeshkar Hospital, Pune
- BGS Global Bangalore
- GCRI Ahmedabad
- Rajiv Gandhi Cancer Institute, Delhi
- Apollo Hospital, Ahmedabad
- AIMS, Kochi
- Sahyadri Superspeciality Hospital
- Manipal Hospitals, Bangalore
- Sterling Hospital, Ahmedabad
- ARTEMIS Hospital, Gurgaon
- MIOT Hospital, Chennai
- Apollo Hospital, Delhi
- Meenakshi Mission Hospital & Research Center
- Apollo Glen Eagles Hospital, kolkata
- Basavatarakam Indo American Cancer Hospital, Hyderabad
- Cancer Institute, Adyar
- Christian Medical College (CMC), Ludhiana
- HCG, Bangalore
- PD Hinduja Hospital, Mumbai
- SRMC, Chennai
- Tata Medical Center, Mumbai

Reference:

http://datri.org/transplant-centers/



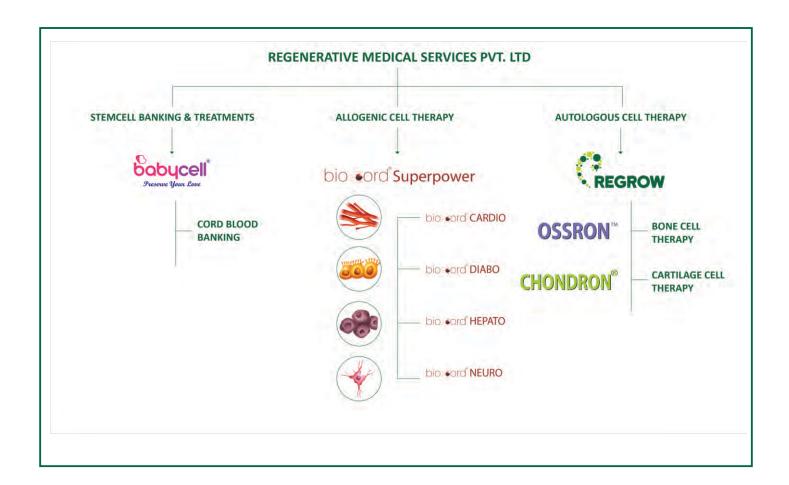


About us

Regenerative Medical Services Pvt. Ltd (RMS) started its journey in 2009, when the concept of umbilical cord blood banking in India was just another science experiment; the outcome of which was unknown to most.

Babycell[™], established in 2009, is India's premium Cord Blood Bank. We understand your baby means everything to you, hence we strive to preserve your love in the safest and purest environment possible. We are leaders not only in umbilical cord blood preservation but our value-added services expand in providing Diagnosis, best Processing Technology and customized Therapeutic Solutions for you and your loved ones.

Driven with passion and powered with cutting edge technology, BabycellTM has become one of the preferred partners for stem call banking in India. BabycellTM has gained the trust of more than 20,000 thousand families in a short span of time and is growing family by family.





Milestones

- Founded in 2009 as the first Indian biotechnology company to introduce cutting edge technology platform for regenerative medicine in India

 Achieved ISO in the very first year of operations
 Appointed Deloitte India, the global leaders in auditing as our financial auditors from the year 2010 till date
 Introduced New Born Screening
 - Selected as one of the top five Leaders of Tomorrow by ET Now

operations

on CNBC Young TurksSelected as one of the top five Leaders of Tomorrow by ET

Satyen Sanghavi, CSO featured as the Young Entrepreneur

- Selected as one of the top five Leaders of Tomorrow by ET Now again!
- Introduced Sickle Cell Screening

Businesses & stars of the Industry

- Product Innovation Award by Asian Confederation of
- Invited for Bio-Africa conference, Mauritius to speak about RMS's advanced regenerative medical therapies
- Satyen Sanghavi, CSO featured on CNBC Africa's Power Lunch, giving an update on stem cell technology
- 1st Cord blood transplantation approval for a Patient diagnosed with Mucopolysaccharidosis Type VI (MPS VI) carried out by Dr. Revathy Raj (Apollo Speciality Hospital, Chennai).
- Cord Blood transplatation was performed for baby suffering from -Thalassemia Major, carried out by Dr. Vijay Ramanan (Ruby Hall Clinic, Pune).
- RMS Laboratory received inhouse R&D unit recognition from Department of Scientific and Industrial Research(DSIR)
- RMS received Market Authorization for Chondron and Ossron along with filing of patent for Chondron, Ossron and Uregrow

 Presented at Stem Cells Asia Regenerative Medicine Congress, Singapore

Certified with GMP, GLP and GCP in its 2nd year of

- Received the Healthcare Excellence Award For Cord Blood Bank by Worldwide Achievers and Headlines Today
- Introduced Mummy & Tummy events for pregnant ladies

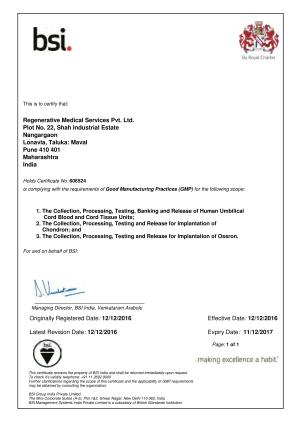
- Launch of Sibling Program as CSR for children requiring stem cell transplant.
- SKOCH order of merit awarded for India's Best SME's (2015)
- Invited to Global Business Summit
- Initiated contribution of a portion of the sales proceeds towards Cancer patients treatment

RMS got DCGI approval to market cell therapy drug Ossron and Chondron
 Exclusively associated with Apollo Hospitals to provide
 Regrow services across India.

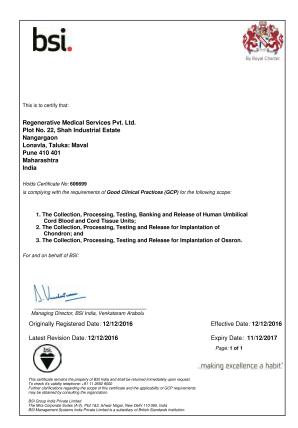




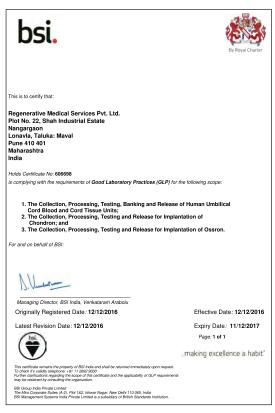
Accreditation



GMP (Good Manufacturing Practices) 2016-17



GCP (Good Clinical Practices) 2016-17



GLP (Good Laboratory Practices)2016-17

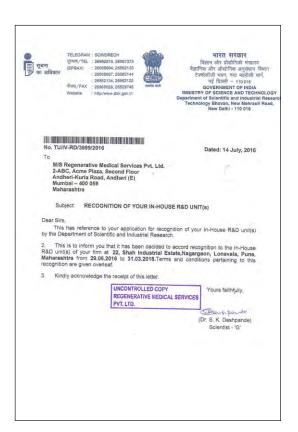




Accreditation



ISO 13485:2003
This is specifically for medical manufacturing 2016-17



Recognition of R & D unit by DSIR 2016-17



Awards



The top 5 Leaders of Tomorrow by ET Now for 2 consecutive years: 2011and 2012



Product Innovation Award by Asian Confederation of Businesses and Stars of the industry (2014)



Healthcare Excellence Award for Cord Blood Bank by Worldwide Achievers (2013)



India's Best SME's (2015) by Skotch Order of Merit.



Recognitions

5th annual





Speaker and Presenter at Stem Cells Asia Regenerative Medical Congress, Singapore

Speaker at
Bio Africa Conference,
Mauritius







Featured in CNBC Young Turks



Selected as Top 5 Leaders of Tomorrow by ET Now



Cell Processing Centre[™]



























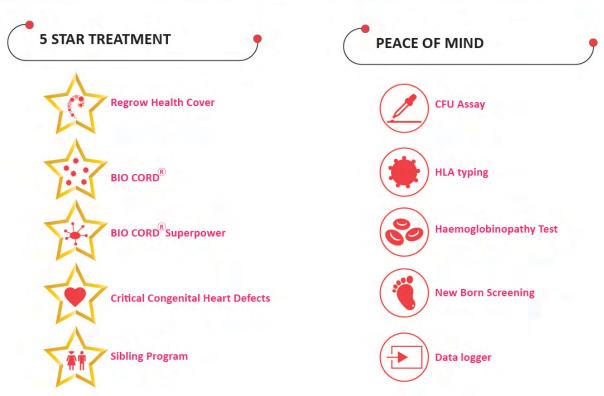




Babycell Presence Maur. Laue

Why Babycell™







Saving Cord Blood is easier than you think



■ Meet

Our key account manager will explain various benefits and guide you through the enrolment process

Receive ▶

Upon registration, Babycell™ collection kit will be sent to you



24

◄ Collect

Call 1800-209-0309 on your way to the hospital. Bring your kit along for sample collection

Transport ►

Logistics team will safely transfer the sample to Babycell[™] Cell Processing Centre





◄ Preserve

With the Certificate of Cryopreservation you can rest assured your baby's health is preserved for life



Transport specialists will access the medical condition and prescribe suitable transplantation procedure







Cryopreservation Report



Client	ID:		Lab ID :	
Mothe	r's Name (Client) :		Father's Name :	
Date C	Of Collection:		Date Of Report :	
Date 0	Of Processing:	- 1		
Produ	et: Cord Blood			
Natur	e Of Packing and Content	Cells (20 ml pro	Blood Unit Containing Her occessed CB With 5 ml cry overed with overwrap bag	oportectant) Stored in
Stora	ge Condition	Product stored	in Vapour phase of Liquid	d Nitrogen
Sr No.	Test	s	pecification	Results
1	Blood Group	Blood Group (AE	3O) And Rh Factor	
2	Total Nucleated Cell Count Automated Cell Counter	NLT 50 X 10 • /	unit	
3	Total Mononucleated Cell Count Automated Cell Counter	Observed Value	/ unit	
4	Total CD 34+ Stem Cell Count By Flow Cytometry	NLT 1.3 X 10 ● / unit		
5	TNCC Cell Viabillity (%) By Flow Cytometry	NLT 70%		
6	CD34+ Cell Viability (%) By Flow Cytometry	NLT 70%		
7	Hematocrit (%)	Observed Value (%)		
8	Sterility Testing* By Direct Inocultaion Method Automated Microbial Detection System	Negative		
NL	Richard Stood Processed and Cryopreserved Suc T: Not Less Than terility is positive please refer microbial iden		d ()	
11 5	termy to positive, predoc refer thiolopial iden	misation and not repo		
				Medical Director
				,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,
				Ref SOP/QA/GEN/023
		vices Pvt.Ltd.22, Shah		A STATE OF THE STATE OF

DSIR V V GAP V V GAP V V GCP V V GLP



Cryopreservation Report



Certificate of Cryopreservation

Client ID:		Lab ID :		
Mother's Name (Client) :		Father's Name :		
	a transfer of the state of the			
Produ	ct : Maternal Blood			
Sr.No	Test	Specification	Results	
1	HIV I & II Antibodies	Negative		
2	HBs Ag (Surface Antigen)	Non-reactive / Reactive		
3	HBc Antibodies	Non-reactive / Reactive		
4	HCV Antibodies	Non-reactive / Reactive		
5	HTLV I & II Antibodies	Negative / Positive		
6	CMV IgM	Negative / Positive		
7	CMV IgG	Immune / Non-Immnue		
8	Malarial Parasite	Not Detected / Detected		
9	Syphilis Antibodies	Non-reactive / Reactive		

Remarks :-		

Medical Director

Dr.Yogesh Gavandi

(M.B.B.S.,M.D.(Pathology))

Ref SOP/QA/GEN/023

1. Cryopreservation report Disclaimer & Confidentiality

Disclaimer: The Laboratory of Regenerative Medical Services Private Limited (RMS Lab) is certified annually to ISO 13485 standards, and regularly inspected by the FDA. Whilst every effort has been made to ensure the accuracy of the information contained in this report, all investigations are carried out with the highest professional standard of due care and skills, based on the scientific knowledge by our medical experts but limited to sensitivity and specificity of individual assay as well as the Umbilical Cord Blood received by the Laboratory. Regenerative Medical Services Pvt. Ltd, its officers, directors, agents, attorneys, employees, shareholders, and affiliates do not take any responsibility or liability for any errors that may happen in the pre-analytic, analytic and post-analytic phase of testing due to failure of the systems, process and procedure which is beyond the control of RMS Lab involved in the transmission and reporting of the test result. Isolated laboratory investigation does not confirm final diagnosis.

Confidentiality: This is a confidential report from Regenerative Medical Services Pvt. Ltd. The contents of this report and any attachments are solely intended for the addressee(s) and in all likelihood, contains information that is legally privileged. If you are not the intended recipient of this report, we request you to kindly return the copy immediately. Please also simultaneously disregard the contents herein and destroy any copies, if any created.

Lab Address : Regenarative Medical Services Pvt.Ltd.22, Shah Industrial Estate,Nagargaon, Lonavala - 410401, Tal:Maval, Dist: Pune,M.S.India. Tel : (+91) 02114-273741-42-43.

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Cryopreservation Report



CFU Report

Client ID:	Lab ID :	
Mother's Name (Client) :	Father's Name :	

Hematopoietic Colony Forming Units Assay (CFU Assay)

Product: Cord Blood			
Sr No.	Test	Specification	Results ***
01	BFU-E*	Growth (for positive results of Potency)	
02	CFU-GM**	Growth (for positive results of Potency)	





Importance of CFU: This test is an indication of the proliferative capacity of the stem cell, which reflects upon the regenarative capacity of the Cryopreserved hematopoietic stem cells.

Remarks :-

* BFU-E: Burst Forming Unit-Erythrocyte

** CFU-GM : Colony Forming Unit-Granulocyte Macrophage

*** Value is representative of total hematopoietic progenitor colonies in 25 ml cryopreserved cord blood

Date: 12.06.2017 Medical Director

Ref SOP/QA/GEN/023

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HLA Typing Certificate

Testing partner: Histogenetics

HISTOGENETICS

300 Executive Blvd, Ossining, New York - 10562.
Phone: 914-762-0300. Fax: 914-762-4441. Website: www.histogenetics.com
ASHI # 03-1-NY-26-2. CLIA # 33D0985173
Soo Young Yang Ph.D / Nezih Cereb M.D., Directors
Email: customerservice@histogenetics.com

HLA TYPING REPORT

Sample Information Last Name Client ID Sample ID Collected On : Received On : DRB1* DRB345* DQB1* DPB1* DPA1* 02:11:01G 14:04:01 40:06:01G NA NA NA NA NA NA 24:02:01G 40:01:01G NA 15:02:01G NA NA lΝΑ NA

Typing Status : Complete

Null Allele Resolution Status :

Note:

- 1. Allele Database Version: 3.25.0 (Jul 2016)
- 2. Allele bearing suffix G: The first allele of the included Alleles. Included alleles in the appendix
- 3. Sequencing Technology: Illumina Miseq
- 4. Reagent Information: Lot Number available upon request.

Report History	Report History				
Reported Date		Comments			
29-Jun-2017					

DIRECTOR / ASSOCIATE DIRECTOR

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Page 1 of





HLA Typing Certificate



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Soo Young Yang Ph. D / Nezih Cereb M.D., Directors
Email: customerservice@histogenetics.com

HLA TYPING REPORT - APPENDIX

Locus	G Code	NMDP Allele Code	Exon Sequenced	Included Alleles
Sample	e ID :			
Α	02:11:01G	02:AZGG	Exon 2,3	02:11:01/02:69
Α	24:02:01G	24:ABGEW	Exon 2,3	24:02:01:01/24:02:01:02L/24:02:01:03/24:02:01:04/24:02:01:05/
				24:02:01:06/24:02:01:07/24:02:01:08/24:02:03Q/24:02:10/
				24:02:13/24:02:31/24:02:40/24:02:43/24:02:44/24:02:56/
				24:02:65/24:02:79/24:02:80/24:02:81/24:02:82/24:02:83/
				24:02:84/24:09N/24:11N/24:40N/24:76/24:79/24:83N/24:144/
				24:150/24:153/24:154/24:155N/24:163N/24:183N/24:231/24:249/
				24:250/24:251/24:263/24:264/24:265/24:266/24:267/24:268/
				24:269/24:270/24:271
В	40:01:01G	40:ANAGY	Exon 2,3	40:01:01/40:01:02/40:01:25/40:01:36/40:01:37/40:01:42/40:01:45/
				40:55/40:141/40:150/40:151/40:179/40:221/40:236/40:241/40:247/
				40:264/40:272/40:278/40:299/40:301/40:329/40:338N
В	40:06:01G	40:06	Exon 2,3	40:06:01:01/40:06:01:02/40:06:03/40:06:04/40:06:08
DRB1	14:04:01		Exon 2	
DRB1	15:02:01G	15:02:01	Exon 2	15:02:01:01/15:02:01:02

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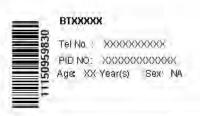
Histogenetics lab is certified by
ASHI - American Society for Histocompatibility & Immunogenetics
CLIA - Clinical Laboratory Improvement Amendments





Heamoglobinopathy Report

Testing partner: Metropolis



Reference

Sample Collected At:

VID:XXXXXXXXXXXXXXIIIected On:

Registered On: XXXXXXXXXXXXXX AM Collected On: XXXXXXXXXXXXXAM Reported On:

XXXXXXXXXXXXXX PM

Abnormal Haemoglobin StudiesHb VariantsBlood

(XXX Whole Blood)

	VIII. 1777-1-1-1-1		
Investigation Foetal Haemoglobin (HbF)	Observed Value	<u>Unit</u> %	
, cara gama gama cany	210		
Haemoglobin A0 (Hb A0)	XX	%	
Haemoglobin A2 (HbA2)	XX	%	
Impression			
	History provided : - Cord blood sample.		
	Kindly correlate with ag	ge of the patient.	
Comment	¥ ***		
Suggestion	2		
Method	HPLC		

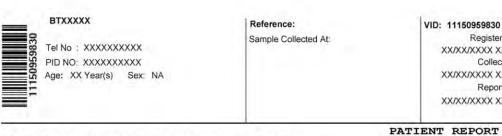
Interpretations & Remarks:-

- All results have to be correlated with age and history of blood transfusion if there is history of blood transfusion in last 3
 months, repeat testing after 3 months from last date of transfusion is recommended.
- In case of haemoglobinopathy, parents or family studies and counseling is advised.
- This test detects Beta thalassaemia and haemoglobinopathies. DNA analysis is recommended to rule out alpha thalassaemia and silent carriers.
- Linearity range of HbF is 1-40%, however, values in excess of the reportable range have been provided for ease of interpretation.
- Mild to moderate increase in fetal heamoglobin can be seen in some acquired conditions like Pregnancy, Megaloblastic anaemia, Thyrotoxicosis, Hypoxia, Chronic kidney disease, Recovering marrow, MDS, Aplastic anaemia, PNH, Medications (Hydroxyurea, Erythropoletin) etc.
- P3 window- Above 10% is often indicative of either denatured forms of hemoglobins or may suggest a possibility of abnormal haemoglobin variant. Hence repeat analysis with fresh sample or DNA studies is advised.
- P2 window- Above 10% is indicative of either glycated haemoglobin requiring correlation with diabetic status or may suggest
 a possibility of abnormal haemoglobin variant requiring further DNA studies for confirmation.





Heamoglobinopathy Report



Bio-Rad Variant-II, Serial # 12853

BTXXXXX

PATIENT REPORT V2_BThal

Registered On:

Collected On: XX/XX/XXXX XX:XX AM

Reported On: XX/XX/XXXX XX:XX PM

XX/XX/XXXX XX:XX AM

Patient	Data
Sample :	ID:
Patient	ID:

Physician: Sex: DOB: Comments:

11150959830

Analysis Data Analysis Performed: Injection Number: Run Number: Rack ID: Tube Number: Report Generated: Operator ID:

XX/XX/XXXX XX:XX:XX 732

XX/XX/XXXX XX:XX:XX

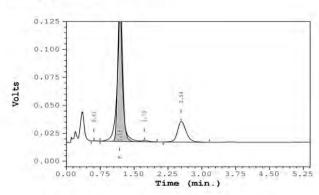
Peak Name	Calibrated Area %	Area %	Retention Time (min)	Peak Area
Unknown		0.1	0.61	1299
F'	80.1*		1.18	870365
P3	(4)4(4)	1.1	1.73	12793
Ao	- are	21.8	2.54	245849

Total Area: 1,130,307

80.1* % F Concentration = A2 Concentration =

"Values outside of expected ranges

Analysis comments:



-- End of Report --

Dr. XX MD(Path) Page 3 of 3

Metropolis lab is certified by NABL - National Accreditation Board for Testing & Calibration Laborities **CAP - College of American Pathologists**





New Born Screening

Testing partner: Nextline Diagnostics (Bharat serum)

NBS list from dried blood specimen

A. Inborn Errors of Amino Acid Metabolism

- 1 Phenylketonuria (PKU)
- 2 Benign hyperphenylalaninemia
- 3 Defects of biopterin cofactor biosynthesis
- 4 Defects of biopterin cofactor regeneration
- 5 Maple syrup urine disease
- 6 Classical citrullinemia
- 7 Citrullinemia type-2 (citrin deficiency)
- 8 Arginosuccinate synthase (ASS) deficiency
- 9 Arginosuccinic aciduria
- 10 Hyperornithinemia, Hyperammoninemia Hyperhomocitrullinemia (HHH) syndrome
- 11 Arginase deficiency/Argininemia
- 12 Neonatal Tyrosinemia
- 13 Tyrosinemia I
- 14 Tyrosinemia type II
- 15 Tyrosinemia III
- 16 Homocystinuria
- 17 Hypermethioninemia

B. Inborn Errors of Fatty Acid Metabolism

- 18 Carnitine transporter deficiency (Carnitine uptakedeficiency)
- 19 Carnitine/acylcarnitine Translocase deficiency
- 20 Carnitine palmityl transferase deficiency type I
- 21 Carnitine palmityl transferase deficiency type II
- 22 Short chain acyl co A dehydrogenase deficiency
- 23 Short chain hydroxy acyl co A dehydrogenase deficiency
- 24 Medium chain acyl CoA dehydrogenase deficiency
- 25 Medium chain ketoacyl coA thiolase deficiency
- 26 Very long chain acyl-CoA dehydrogenase deficiency
- 27 Long chain L-3-hydroxy acyl-CoA dehydrogenase deficiency

- 28 Trifunctional protein deficiency
- 29 Multiple acyl-CoA dehydrogenase deficiency/Glutaric acidemia type II
- 30 Medium chain L-3 hydroxy acyl coA dehydrogenase deficiency

C. Inborn Errors of Organic Acid Metabolism

- 31 Glutaric acidemia type I
- 32 Isovaleric acidemia
- 33 2-Methylbutyryl-CoA-dehydrogenase deficiency
- 34 Propionic acidemia
- 35 Methylmalonyl-CoA mutase deficiency
- 36 Methylmalonic acidemia
- 37 Methylmalonic acidemia (mutase)
- 38 3-Methylcrotonyl-CoA carboxylase deficiency
- 39 Malonic aciduria
- 40 3-Hydroxy 3-methyl glutaric acidemia
- 41 β-Ketothiolase deficiency
- 42 Multiple CoA carboxylase deficiency
- 43 Isobutaryl CoA dehydrogenase deficiency
- 44 Malonyl-CoA decarboxylase deficiency
- 45 Holocarboxylase deficiency
- 46 3- Methyl glutaconyl Co A hydratase deficiency

D. Biochemical DELFIA TRF

- 47 Congenital Hypothyroidism
- 48 Congenital Adrenal Hyperplasia
- 49 Galactosemia
- 50 Glucose-6-Phosphate Dehydrogenase Deficiency
- 51 Cystic Fibrosis
- 52 Biotinidase Deficiency

Biological reference intervals

Analyte	(μM)
C0-carnitine	12 - 100
C2-carnitine	< 72
C3-carnitine	< 6.34
C4-carnitine	< 1.07
C5-carnitine	< 0.48
C5DC-carnitine	< 0.56
C6-carnitine	< 0.17
C8-carnitine	< 0.24
C10-carnitine	< 0.35
C12-carnitine	< 0.41
C14-carnitine	< 0.50
C16-carnitine	< 9.89
C18-carnitine	< 2.06

Analyte	(μM)
Alanine	< 736
Arginine	< 55
Aspartic acid	< 420
Citrulline	< 50
Glutamic acid	< 1073
Glycine	< 1001
Leucine/Isoleucine	< 300
Methionine	< 44
Ornithine	< 453
Phenyalanine	<120
Tyrosine	< 248
Valine	< 212





Mummy & Tummy Program

Babycell proudly presents 'Mummy Ki Paathshaala', where all the lovely to be mommies get a chance to experience going back to school and learning about motherhood!

Mummy ki Paathsaala is an interactive series of sessions to highlight information about pregnancy diet, pregnancy yoga and exercises, labor and delivery tips

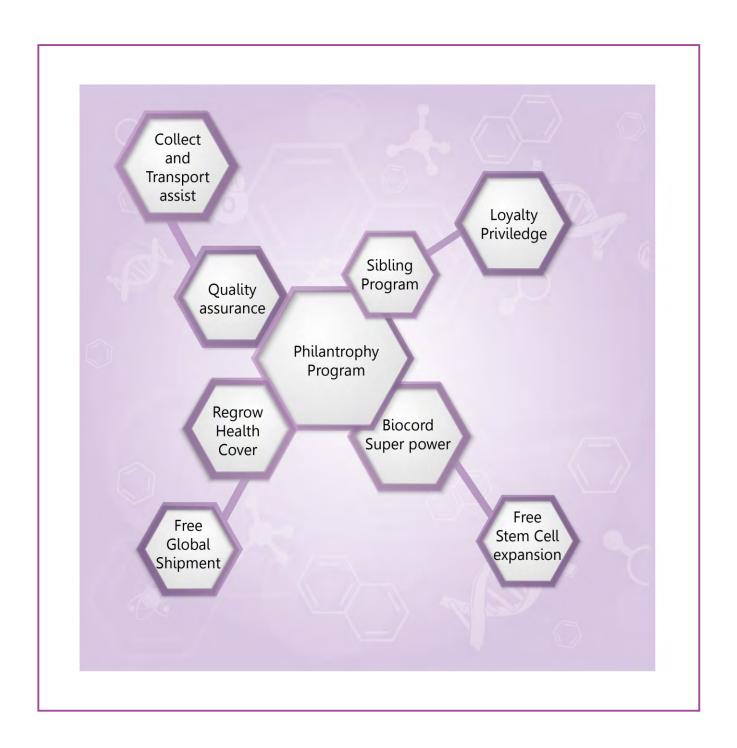
'Mummy Ki Paathshaala' not only helps answer your questions about child delivery and antenatal care, but also helps you find your way to being an awesome parent!

MKP covers some important aspects of pregnancy like the following:





Client Benefit Program





Regrow Therapies



India's first FDA approved permanent, easy and natural Cartilage cell therapy



BENEFITS

- Targeted, personalized Cell Therapy
- Regeneration of hyaline like cartilage
- Minimally invasive procedure
- Further progression of osteoarthritis is ceased
- Complete restoration of range of motion
- Active normal life involving sports

DON'T REPLACE, REGROW.

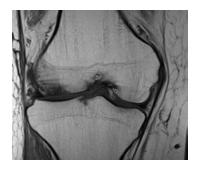
Regrow your Cells. Rebuild your Life^R.



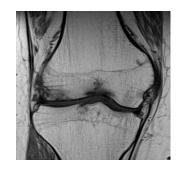


Case Studies

1 Indication : Grade IV Osteochondritis Dissecans Lateral Femoral Condyle Left Knee

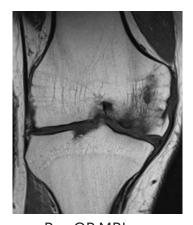




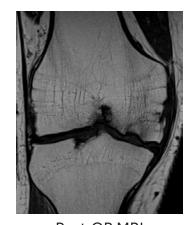


Post-OP MRI

2 Indication: Grade IV Osteochondral defect Medial Femoral Condyle Right Knee







Post-OP MRI

3 Indication : Grade IV Osteochondral defect Medial Femoral Condyle Left Knee



Pre-OP MRI



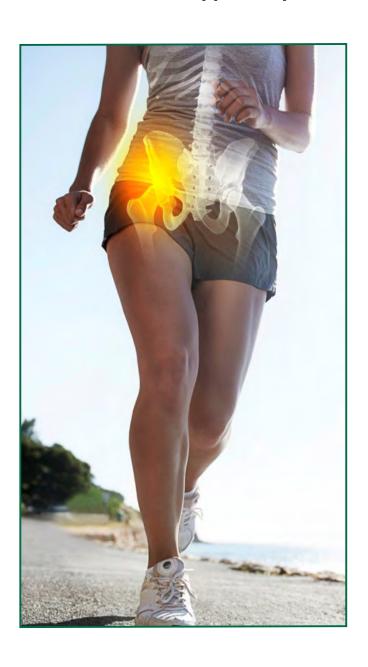
Post-OP MRI



Regrow Therapies

OSSRON[™] Bone Cell Therapy

World's first FDA approved permanent, easy and natural bone cell therapy



BENEFITS

- Targeted, personalized Cell Therapy
- Accelerated bone healing
- Three-dimensional bone regeneration
- Complete relief from pain
- New blood supply is initiated which further supports bone growth
- Active normal life

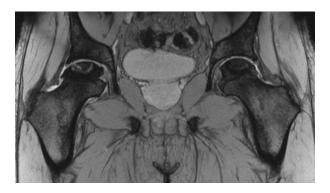
DON'T REPLACE, REGROW.Regrow your Cells. Rebuild your Life[®].

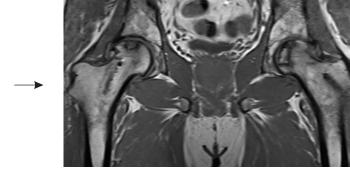




Case Studies

1 Indication: Left unilateral AVN Grade: II

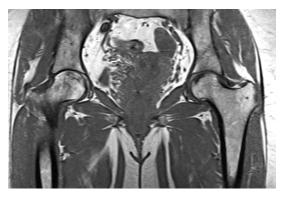




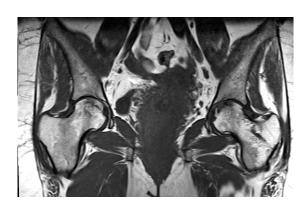
Pre-OP MRI

Post-OP MRI

2 Indication: Right Unilateral AVN Grade: II

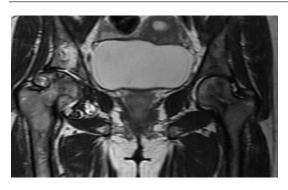




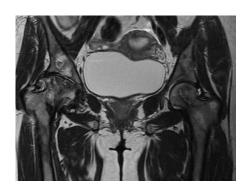


Post-OP MRI

3 Indication: Left Unilateral AVN Grade: II



Pre-OP MRI



Post-OP MRI



Spreading Smiles

Cord blood Stem cells saved Prerna from Dreadful Diseases MPS6

At the age of 2, Prerna's parents sensed something wrong with her structural formation and hence took doctors' opinions. Post a thorough check from specialist of Hinduja hospital, Mumbai. Prerna was diagnosed with Mucopolysaccharidosis Type VI popularly known as MPS 6. It is caused by deficiency of the enzyme N-acetyl- galactosamine-4-sulphatase. The symptoms for this disease are weak bone, bone cartilage disorders, enlargement of liver and spleen. On doctor's suggestion, Prerna's parents went for another child and stored their second child's umbilical cord blood for Prerna's treatment of Mucopolysaccharidosis Type VI metabolic condition (MPS-VI). Prerna's parents approached Babycell under sibling program. On 19th January 2016, the transplant took place at Apollo Specialty Hospital, Chennai by Dr. Revathi Raj. The transplant was successful and Prerna has since then has shown tremendous improvements and after 1 year of follow up, we learnt that Prerna is recovering well.

Disclaimer: Names of the patient have been changed to protect the identity of the patient and family.





Cord blood Stem cells saved Veena from Thalassemia Major

Veena was born with B- Thalassemia Major. Thalassemia major occurs when a child inherits two mutated genes one from each parent. The symptoms of these disease are fatigue, weakness, jaundice, swollen abdomen, dark urine and deformities of the facial bones. Veena's parents did not store her cord blood at birth. Since Veena had no permanent cure for her Thalassemia Transplant, Physician Dr. Vijay Ramanan suggested Veena's parents to have another baby so that they can use their second child's cord blood stem cells to cure Veena from Thalassemia. Veena's parents approached Babycell for her treatment under their sibling program. On 29th July 2016, the transplant took place at Ruby Hall Clinic, Pune. The transplant went well and after 6 months of follow-up, Veena is seen recovering well. Moreover, there are no signs and symptoms of the disease. Today, the family is relieved as they had, once, lost hopes of her recovery. Disclaimer: Names of the patient have been changed to protect the identity of the patient and family.



Press Releases

Business Line

The Hindu • Business Line Thursday, July 21, 2011

Babycell reports spurt in demand to store umbilical cords

Madfumath D.S.

Bangalora, July 19

As if monsoon discounts and off-season FMCG shavers are noted on the state of banks at the cell banks of the state of of



BusinessLine

Apollo Hospitals ties up with cell therapy firm



HARMABIZ . com

ollo & RMS REGROW bring to India first of its kind innovative cell therapy atment for orthopedic patients

hindustantimes

Apollo Hospitals bring innovative cell therapy to treat joint pain

Apollo Hospitals and RMS REGROW will bring to India first of its kind innovative cell therapy for orthopaedic patients.



Dr Prathap C Reddy, founder-chairman, Apollo Hospitals (2nd from left) announcing the partnership to bring innovative cell therapy to India (HT ph

Making uses of the advances in cell therapy, Apollo Hospitals on Tuesday announced partnership with RMS REGROW, to offer new regenerative medicine cell therapy products that will be a boon for patients suffering from bone and cartilage damage.

The two products- Ossron and Chondron, will address unmet clinical needs in the orthopaedic market with respect to sports injuries, accidents and alternate to hip replacements and knee replacements for a young arthritic knee.

In an exclusive tie up between Apollo Hospitals and RMS Regrow, the treatment therapy will be made available across all Apollo Hospitals, Apollo Spectra Hospitals and Apollo clinics in India.



RECOGNIZED AS RESEARCH & DEVELOPMENT UNIT

by the Department of Science and Industrial Research (DSIR) Ministry of Science and Technology, Govt of India











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CIN: U24100MH1989PTC054162 | GST No. 27AADCS4917N1ZW

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