

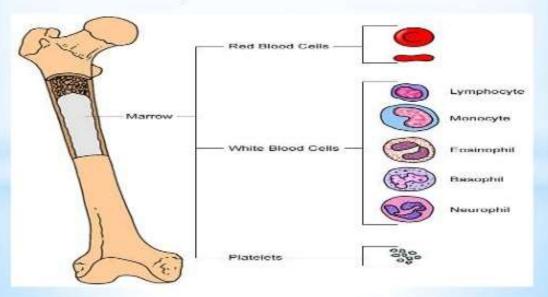


Patient Flow Predictive Model

WHAT IS THE BONE MARROW ?!

Bone Marrow is the soft spongy tissue that fills the cores of larger bones.

It serves an active function in the body by producing all three types of blood cells, as well as lymphocytes, wich support the immune system.





Types of Bone Marrow Transplant

Autologous Transplants

Here stem cells are removed from the patient before highdose chemotherapy or radiation treatment. After chemotherapy your stems cells are put back in your body to make (regenerate)

normal blood cells

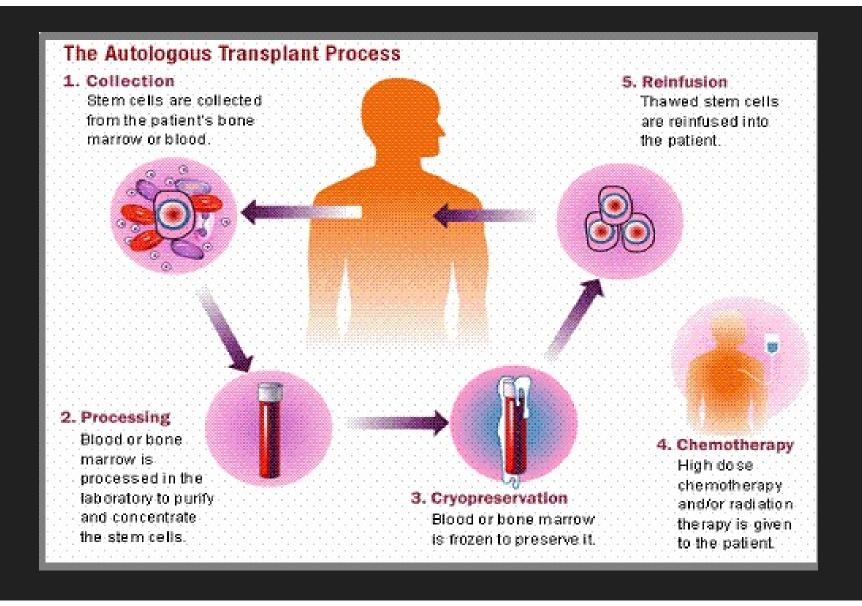
Allogeneic Transplants

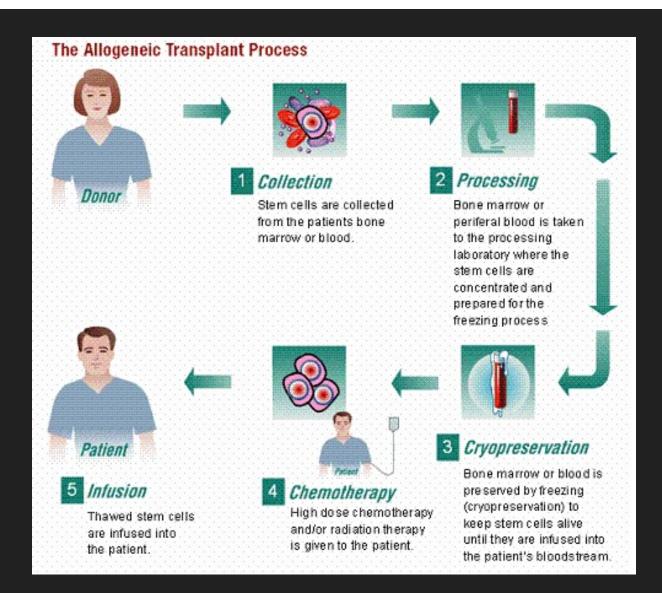
Stem cells are removed from another person, called a donor. Donor's genes must at least partly match your genes, which are decided through special blood tests. Brother, sister, parents, children are most likely to be a good match.

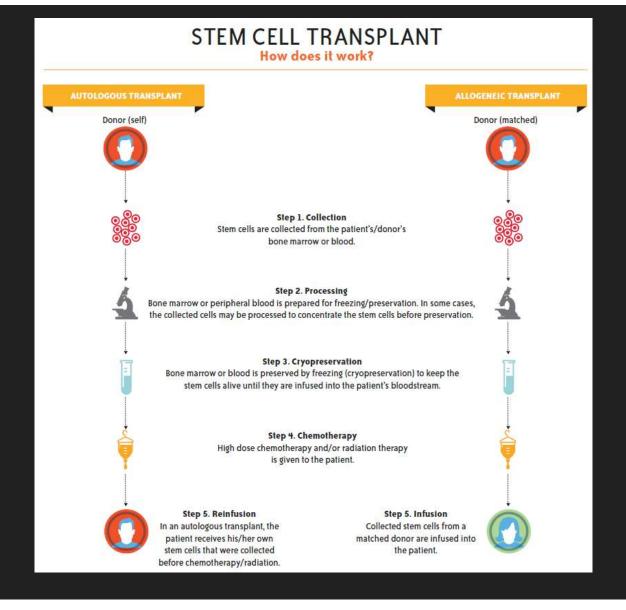
Umbilical cord blood transplant

Stem cells are removed from a newborn baby's umbilical cord right after birth. The stem cells are frozen and stored until they are needed for a transplant. These cells are very immature so there is less of a need for matching.









Donor Search		Conditioning (Chemo and/or radiation)	Cell Infusion "Day 0"	In-Patient Recovery	Out-Patient Recovery	At Home Recovery	The "New Normal"
Adult	Cord blood: About 2 weeks	Standard (Myeloablative) transplant					
donor:		6–12 days	"Day 0"	30–100 days	100 days or more	100 days or more	1 year or more
About 2 months		Reduced Intensity (Non-Myeloablative) transplant					
or longer		Varies	"Day 0"	Varies	Varies	Varies	1 year or more

Project Goal:

Create a predictive model from historical data that will estimate patient volume through the various clinical phases of the Bone Marrow Transplant (BMT) process.

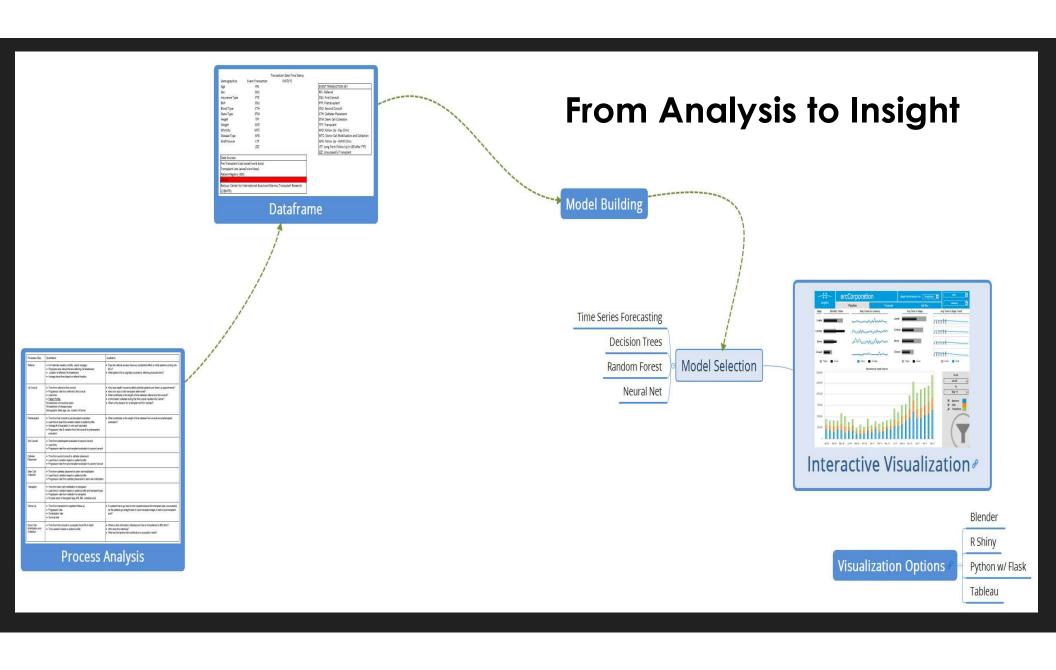
Data from 2010-2015 will be used to train the model.

Data from 2016-2017 will be used to test and validate model accuracy.

Project Outputs:

Dashboard that shows red, yellow or green representation of patient volume through each phase of the process

Estimate of quarterly and annual transplant volumes



Processes Step:	Quantitative	Qualitative
Referral	## of referrals (weekly,monthly, yearly average) Physicians and clinics that are referring (% breakdown) Cocation of referrals (% breakdown) Average travel time based on referral location	Does the referral process have any substantial effect on initial patients coming into MCV? What patient info is originally provided by referring physician/clinic?
1st Consult	Time from referral to first consult Progression rate from referral to first consult Lead time Patient Profile: breakdown of insurance plans breakdown of disease types Demographic Data (age, sex, location of home)	How does health insurance affect potential patients and follow up appointments? How is an auto or allo transplant determined? What contributes to the length of time between referral and first consult? Is information collected during the first consult inputted into Cerner? When is the decision for a transplant at MCV decided?
Pretransplant	Time from first consult to pre-transplant evaluation Lead time & lead time variation based on patient profile Average # of evaluation in one work day/week Progression rate & variation from first consult to pretransplant	What contributes to the length of time between first consult and pretransplant evaluation?
2nd Consult	Time from pretransplant evaluation to second consult Lead time Progression rate from pre transplant evaluation to second consult	
Catheter Placement	Time from second consult to catheter placement Lead time & variation based on patient profile Progression rate from pre transplant evaluation to second consult	
Stem Cell Collection	Time from catheter placement to stem cell mobilization Lead time & variation based on patient profile Progression rate from catheter placement to stem cell mobilization	
Transplant	Time from stem cell mobilization to transplant Lead time & variation based on patient profile and transplant type Progression rate from collection to transplant Moreover by break down of transplant type (PB, BM, umbilical cord)	
Follow Up	Time from transplant to outpatient follow up Progression rate Complication rate Survival rate	 If a patient has to go back to the hospital because the transplant was unsuccessful, do the patients go straight back to input transplant stage or back to pre transplant eval?
Donor Cell Mobilization and Collection	Time from first consult to successful donor/HLA match Time variation based on patient profile	Where is this information collected and how is it transferred to BM clinic? Who does the matching? What are the factors that contribute to a successful match?

The Process

Trans	action	Date	Time	Stamp
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		Transaction Date Time Stamp
Demographics	Event Transaction	(M/D/Y)
Age	RFL	
Sex	CN1	
Insurance Type	PTP	
BMI	CN2	
Blood Type	CTH	The Data
Gene Type	STM	ille Dala
Height	TPT	
Weight	AFD	
Ethnicity	MTC	
Disease Type	AFG	
Graft Source	LTF	
	ZZZ	

Data Sources:

Pre-Transplant Lists (excel/word docs)

Transplant Lists (excel/word docs)

Patient Registry (IDX)

Cerner

Backup: Center for International Bood and Marrow Transplant Research (CIBMTR)

EVENT TRANSACTION KEY

RFL: Referral

CN1: First Consult

PTP: Pretransplant

CN2: Second Consult

CTH: Catheter Placement

STM: Stem Cell Collection

TPT: Transplant

AFD: Follow Up - Day Clinic

MTC: Donor Cell Mobilization and Collection

AFG: Follow Up - GVHD Clinic

LTF: Long Term Follow Up (+100 after TPT)

ZZZ: Unsuccessful Transplant

Next Steps:

- Develop business rules for transactions and date of transaction by data source with retrieval method
- 2) Capture demographic data for each patient
- 3) Capture transaction data for each process
- 4) Join patient and process data via column vector
- 5) Perform analysis and choose best model

Questions/Comments