



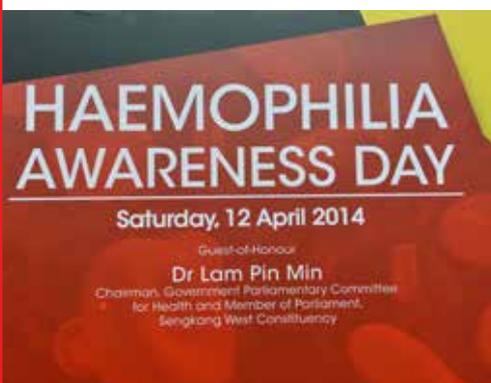
HAEMOPHILIA

SOCIETY OF SINGAPORE



ANNUAL REPORT

2013 - 2014



ABOUT US

The Haemophilia Society of Singapore is a voluntary organization that was set up in 1985 by a group of people with haemophilia, parents, healthcare providers and some interested professionals. It is an Institution of Public Character and an affiliated member of the National Council of Social Service and a member of the World Federation of Haemophilia.

The Society's main service to its members is to provide subsidies for members' treatment costs at three government hospitals. This is to alleviate the high costs that members often incur following bleeding episodes. Apart from financial assistance, the Society offers support to parents to help them cope with a haemophilic child, promotes home treatment and keeps members abreast of developments in the area of haemophilia care.

Our Mission

To ensure that every person with haemophilia receives affordable and safe treatment so as to lead full and productive lives in society.

Our Objectives

- * To help people with haemophilia to achieve their potential.
- * To look after the welfare of people with haemophilia.
- * To assist with treatment costs
- * To promote public awareness and understanding of haemophilia

Registered Address

Block 704 Bedok Reservoir Road
#01-3622
Singapore 470704

Mailing Address

Farrer Road P O Box 0273
Singapore 912810

Key Bankers

DBS Bank
Maybank

Auditors

TKNP International

Charity Registration No: 00374
Unique Entity No: S85SS0047A
Website: www.haemophilia.org.sg

Society Registration No: ROS 105/85
For enquiries: 62358166 (Dr Tan HH)
64490289 (Dr Gan KL)

Treatment Centres

Singapore General Hospital Haematology

Staff-in-charge : Lillie Ho
Telephone No. : 63213844

National University Hospital Paediatrics

Doctor-in-charge : Dr Koh Pei Lin
Telephone No. : 67725030

KK Women's & Children's Hospital Paediatric Haematology

Staff-in-charge : Lim Chiew Ying
Telephone No. : 63926343

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PRESIDENT'S MESSAGE



The year started on a promising note. The Society received the “good news” in July 2013 that it was a beneficiary for a fund raising event organized by the students of Hillgrove Secondary School. To receive such support from the community was indeed great

encouragement for us in our quest to better the lives of the members with haemophilia under the Haemophilia Society umbrella. Treatment for haemophilia is expensive and the Society's income is expended mainly on subsidies for treatment costs of its members.

Our Society's focus is to have an effective long term plan for the management of haemophilia needs with co-operation from persons with haemophilia, healthcare professionals and the Ministry of Health. We have a dedicated team of healthcare professionals in the Treatment Centres at Singapore General Hospital, National University Hospital and KK Women & Children's Hospital who provide optimal care for patients with haemophilia. This has enabled Singapore to be ranked alongside developed societies.

Haemophilia, as opposed to other chronic genetic illness, has the potential of a normal outcome, with adequate therapy. This is shown in countries like Sweden, Netherlands and Australia. We hope to see greater use of prophylaxis in children as it will be more cost effective in the long term. The Society hopes to bring such issues to the attention of the authorities and to voice its views on healthcare policies that may directly impact those with haemophilia.

Our recent participation at the May WFH 2014 World Congress held in Melbourne,

Australia, provided opportunities for exchange with healthcare professionals and patients on challenges and solutions to improve treatment, to share insights on patient healthcare issues and care and to network with people around the world. It was indeed an enriching experience for the few of us who were there.

The Society's total expenditure was \$170,339.00 and our total income for the year 2013/14 was \$224,975.00, 153% higher than the previous year, giving us a modest surplus of \$54,636.00. The increase in income was largely attributed to the generosity of our well-wishers and a successful charity film premiere that was held in May 2014. As a small charity, we face many challenges especially in the area of fund raising, but with passion and zeal, we can overcome the odds and achieve! My sincere thanks to Member of Parliament, Ms Low Yen Ling, Chua Chu Kang GRC and who is also the Parliamentary Secretary of Culture, Community and Youth, Baxter Pharmaceuticals (Asia) Pte Ltd, Lee Foundation and Singapore Island Country Club, among others, for their support which helped us to achieve our target. In conclusion, I wish to express my heartfelt appreciation and thanks to the Executive Committee for their dedication and valuable contributions to the work of our Society.

I hope that more members will “volunteer” themselves to continue the work of the Society. I believe that the baton must be passed on to the “younger generation” to helm the Society.

Lastly to our donors, volunteers and well-wishers, thank you for supporting our cause this past year. We look forward to your support in the coming years. Together we can strive for an even better tomorrow.

Dr Tan Hooi Hwa

President

Haemophilia Society of Singapore

HIGHLIGHTS OF THE YEAR

In June 2013, the Haemophilia Society of Singapore was informed by Member of Parliament, Ms Low Yen Ling from the Chua Chu Kang GRC, that it was one of two beneficiaries to benefit from the fund raising efforts of the students from Hillgrove Secondary School. The students were involved in a community project to collect old newspapers from households in the school's vicinity and then sell the newspapers to raise funds for the charities concerned.



MP Low Yen Ling and students visiting residents to collect old newspapers

Group photo of students of Hillgrove Sec School



Cheque Presentation

On 30 July, 2013, Dr Tan Hooi Hwa, Ms See Ek May and Ng Kheng Chew attended the cheque presentation ceremony at the school. The Society thanked MP Ms Low Yen Ling, the Principal, staff and students of Hillgrove Secondary School for their combined effort to extend a helping hand to the less fortunate in the community. The Society was given an opportunity to address the school and acquainted the students about haemophilia, a condition unfamiliar to them. They also met Kheng Chew, a student with this condition.



Haemophilia Awareness Day
"Know My Bleeding Points"

In April 2014, to mark World Haemophilia Day which falls on 17 April, the SingHealth cluster of hospitals, namely Singapore General Hospital and KK Children's Hospital, together with Baxter Healthcare (Asia) and Novo Nordis, involved the Haemophilia Society of Singapore to be a part of the "Haemophilia Awareness Day" event at United Square on 12 April, 2014. The Guest-of-Honour was Dr Lam Pin Min, Chairman Government Parliamentary Committee for Health and Member of Parliament, Sengkang West Constituency. Members of the public learnt more about bleeding disorders from staff present and the various posters, pamphlets and exhibits. The publicity generated created a greater awareness about haemophilia and the existence of the Society and the work it does.



Charity Film Premiere Group Photo :
Drs Tan & Gan, MP Low Yen Ling,
Mr & Mrs Shankar Kaul & Grace Khoo

On 4 May, 2014, the 3D Film "The Amazing Spiderman 2" was screened at Cathay Theatre – this being the Haemophilia Society's first Charity Film Premiere, to raise funds for the Society. The event was sponsored by Baxter Healthcare (Asia). Member of Parliament, Ms Low Yen Ling, who is also the Parliamentary Secretary of Culture, Community and Youth, graced the occasion.

Media publicity was by way of interviews with two members of the Society. Mr Chong Kek Ngow appeared on Channel News Asia regarding medical and costs of living. Six year old Lee Jue Ming and his father were in a special feature by Ministry of Health on how Medifund has helped with Jue Ming's healthcare expenses. Straits Times also

had an article on Jue Ming on 21 April, 2014 and Singapore Press Holdings kindly arranged for a video that showed little Jue Ming at home, at play and in hospital for his infusions.

This was available on-line too. The media effort allowed for greater coverage in terms of wider readership and viewership and enabled the Society to gain from the exposure.

The highlights of the year culminated in the WFH 2014 World Congress which was held in Melbourne, Australia from 11 -15 May, 2014. Dr Tan Hooi Hwa, Dr Gan Kim Loon, See Ek May and Grace Vaz represented the Haemophilia Society of Singapore at the Congress. Their participation was made possible through the kind sponsorships from Baxter Healthcare (Asia) and Biotest. The WFH Congress is considered the largest international meeting for the global bleeding disorders community. The team from Singapore gained much knowledge in terms of advances in treatment and best practice care.

KEY PROGRAMMES

There are about 200 people in Singapore with haemophilia – all males. Haemophilia is usually inherited, but one-third of people have no previous history.

Members of the Haemophilia Society of Singapore comprise babies, toddlers, children, youths, young and mature adults. Growing up with haemophilia involves the whole family. With today's medical advances, parents need not feel it is "the end of the world" when their sons are diagnosed with haemophilia. All these boys can have a bright future once the "missing factor" is replaced.

Effective treatment for haemophilia is available, but as yet there is no cure. Treatment is very costly for those with haemophilia and most working individuals and families would have difficulties meeting treatment costs, especially if they experience frequent bleeding episodes into muscles and joints, which if not managed properly, can cause arthritis and joint problems. Aid from the Society is rendered to all its members regardless of financial background, to help alleviate the heavy costs of treatment.

The key programmes offered by the Society are aimed at addressing the different needs of our members.



- **Treatment Subsidy**
Haemophilia is treated by replacing the missing clotting factor in the blood. This is done by injecting a product that contains the needed factor into the vein. Bleeding stops when enough clotting factor reaches the affected area. The Society assists its members with monthly

subsidies to defray part of their treatment costs, capped at \$800.00 and \$400.00 per month for youths /adults and children respectively. The subsidies' programme is the main expenditure item for the Society. Hence all its funds raised are used mainly to support this programme.

- **Welfare Fund**

The Society has in place a welfare fund, with funds specifically designated to provide financial assistance to members with haemophilia facing financial hardships. Members who require financial aid would need to submit their request and aid is approved when the situation warrants it. The approved applicants are reviewed every quarterly and they are encouraged to seek employment where possible, rather than be dependent on the Society.

- **Educational Awards**

The Society initiated the Educational Awards programme, also known as "Merit Awards" in the year 2000. The awards were to encourage our members in primary, secondary, junior colleges and polytechnics to value the importance of education, that with education they can improve their lives and reach higher heights.

The awards given at each Annual General Meeting are based on academic performance and the two categories are "excellence" and "encouragement" and the monetary rewards range from \$400.00 (junior college/polytechnic) to \$100.00 (primary). This year, 8 awards were given to members from primary (1), secondary (5), Polytechnic (1) and SIM Diploma (1).

The Dulcie Wilson Scholarship Fund in memory of the late Mrs Dulcie Wilson was set up in 2010. A sum of \$20,000.00 was set aside for this purpose. This fund would be used to assist those pursuing studies in a recognized local tertiary institution. Grants of a maximum of \$1,000.00 per academic year would be granted to a maximum of 3 members per year. The scholarships are given on a yearly basis. No applications have been received to date.



Merit Awards' Presentation to Benjamin Tang

- **Seminars / Talks**

Seminars and talks are organized for members of the Society to impart knowledge of haemophilia, to keep members abreast of latest developments and management of haemophilia from specialists in the field, both locally and from overseas. Talks for members range from caring for joints, importance of dental care, to physical activities members can engage in.

- **Support Group**

The Mothers' Support Group which was started in 2009 had been well received. Its purpose was to enable mothers with haemophilic sons to come together to share their experiences in managing their sons' conditions and to obtain moral and emotional support from one another. Most mothers gave positive feedback of the support group as the mothers feel connected as they share a common bond. At these gatherings, mothers and their sons are encouraged to learn self-infusion so that the clotting factor can be administered promptly to minimize bleeding episodes. This would enable them to be on home treatment with minimal disruption to their daily routine.

- **Activities / Outings**

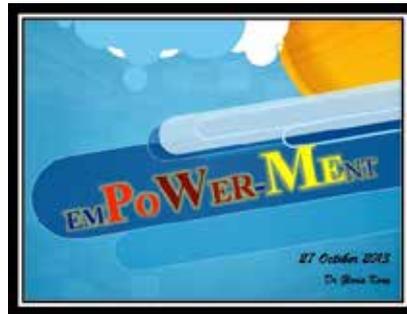
During the year, the Society arranged for different activities or outings for its members. In 2013, some of our members attended two talks by Dr Joyce Lam, Consultant, Paediatric Haematology, KKCH who spoke on "Know Your Factors" and Dr Scott Dunkley, Specialist Haematologist and Director, Haemophilia Treatment Centre at the Royal Prince Alfred Hospital, Sydney, whose topic was "Prophylaxis – Is this still relevant?" In October, members also had the opportunity to hear educational psychologist, Dr Gloria Kong's talk on "EmPowerMent" at the Medical Alumni Theatre. The talk was especially beneficial to the youths who were asked to examine elements within themselves that could help them be the best that they can be.

In 2014, the Society organized outings for members to Sungei Buloh Wetland Reserves, the Nyea Phoa Flower Garden and the Live Turtle & Tortoise Museum. For the Sungei Buloh outing, the Society was assisted by the Nature Society of Singapore, who provided a bi-lingual volunteer to conduct the guided trek of the Reserves.

On 13 April, 2014, to commemorate World Haemophilia Day, more than 50 members of the Society had a sumptuous buffet lunch at the FernTree Café, Miramar Hotel. Food and fellowship made the event memorable!



Dr Scott Dunkley's talk



Slide on "EmPowerMent"



Sungei Buloh Outing



World Haemophilia Day Lunch photo

EXECUTIVE COMMITTEE 2013/14

<i>Name</i>	<i>Designation</i>	<i>Date of Appointment</i>
Dr Tan Hooi Hwa	President	29 September, 2013
Dr Gan Kim Loon	Vice-President	29 September, 2013
Wee Ai Choo	Honorary Secretary	29 September, 2013
Chang Yew Wye David	Assistant Honorary Secretary	29 September, 2013
Lim Keok Kung Freddy	Honorary Treasurer	29 September, 2013

Members:

Ng Teck Hiang	29 September, 2013
Poh Soon Leong	29 September, 2013
Wee Lester	29 September, 2013
See Ek May	29 September, 2013
Ang Har Boon Anthony	29 September, 2013

Haemophilia Society of Singapore is governed by the Executive Committee which has the authority and responsibility for policy making and determination of activities. Members of the Committee are volunteers and receive no monetary remuneration for their contribution.

MY “WORLD CONGRESS” EXPERIENCE

- a parent's perspective

I had the privilege of attending the WFH 2014 World Congress in Melbourne, Australia in May. It was an eye-opener. Never have I met so many people interested in haemophilia. There were possibly a few thousand at the World Congress and they came from all around the world. I met people from the Philippines, Malaysia, Brazil, America, Canada, India, Iran, Australia and many other countries.

Many of those I met are themselves suffering from haemophilia. They are successful people in life, among them are doctors, psychologists, therapists, teachers and businessmen. Haemophilia is just a small disorder in their lives. While some of them suffered from the physical effects of poorly treated injuries, they all worked around their affliction and live successful, fruitful lives. As one of them said, haemophilia is just 5% of their lives. 95% of the time, they are normal and should be treated as normal. I agree totally! I believe that we must look at the positive side and focus on the ability, not the disability.

I was very much inspired by the young adults and teenagers who involved themselves in the Congress, sitting in the sessions, learning more about the disorder and contributing to discussions. It was good to see them interested and taking personal responsibility to do more for themselves and their community. I was touched to see family members, not just parents, but also siblings and wives, who also attended the Congress to support their son/brother/husband in coping with haemophilia.

The Congress catered to a wide range of interests – from medical/dental, to nursing, to psychosocial and musculoskeletal. For me, I attended many workshops and one of the topics that struck me was the one about the impact of haemophilia on family members. I realized that many times we fail to consider the emotional needs of siblings. We should remember not to place the needs of the child with haemophilia above others. How we parent our children and cope with the disorder may impact sibling's relationship with their brother with haemophilia.

Another workshop that resonated with me was one where the speaker reminded the audience not to pamper our children with haemophilia. As with normal kids, pampering can do much harm.

A winning combination for success in life is: “Grit and Good Fortune”. Grit because the individual's passion and perseverance are important and good fortune where one has good doctors, nurses, family and community who can support the treatment and or create opportunities for growth.

My experience at the Congress has shown how important it is for us to come forward to do something for our children, for our community and that all of us have a role to play and to contribute. Let us not wait for others to advocate for us. Instead we need to work together for our children and all who suffer from haemophilia. Only then can we make things happen.



Grace Vaz

JOURNEY WITH HAEMOPHILIA (I)

“ Two of our boys, six year old Lee Jue Ming and fourteen year old Ng Kheng Chew were featured in local publications. We share their journey with haemophilia. ”

LEE JUE MING

– “ The boy who can’t stop bleeding ”

Whenever Lee Jue Ming goes out to play, his mother’s heart skips a beat. This six year old has severe haemophilia, a rare condition that prevents his blood from clotting. He has less than 1 per cent of the normal amount of clotting factors in his blood. He bruises easily and often suffers spontaneous bleeding in the muscles and joints even when there is no injury. If he falls and cuts himself, the result can be devastating. He will need immediate infusion of blood-clotting proteins, also known as clotting factors, to stop the bleeding.



His mother, Madam Ong Lee May, a human resource assistant, would often take Jue Ming and his sister, Wan Ting, 10, to the indoor playground at Changi Airport Terminal One. The playground’s padded surfaces help cushion falls. “The admission is free. We always try to be the first one there at 8 am to avoid the crowd. We would play with him, climb with him and just follow him around in case we need to catch hold of him if he falls,” said Madam Ong.



Jue Ming’s treatment costs of \$1,440 a month are now fully covered by Medifund. This is a relief to Madam Ong, whose take home pay is about \$2000 a month. Her husband, Mr Lee Kwong Luen, quit his job as a mechanic when Jue Ming was six months old to care for him full- time.

“We hope that Jue Ming stays healthy, safe, happy always and can take care of himself eventually. If he can do well in his studies, it will be a bonus,” said Madam Ong.

** This article appeared in the Straits Times on 21 April, 2014*



JOURNEY WITH HAEMOPHILIA (II)

NG KHENG CHEW

– “ Sidelined but not forgotten ”



While other kids run around and play without a care in the world, K.C. cuts a lonesome figure during Physical education lessons, watching from the sidelines.

He suffers from haemophilia and so has to avoid physical activities that might cause bleeding – especially contact sports such as rugby, football and basketball. “When I was younger, I had to stop myself from playing rough, as I will get injured,” says K.C.

“I am most afraid of joint bleeds as they are really painful. These bleeds take a long time to heal and damage my joints. Over time, the joints that suffered bleeding will become ‘targets’, so bleeds will frequently take place at these sites.”

There are the challenges of looking after a person with haemophilia. When he was little, his parents had to watch him round the clock and constantly check his body for signs of bleeding. Like all parents of children with haemophilia, they had to rush off to the hospital as often as four times a week for factor infusion and often meant missing work in the morning.



K.C. needs regular intravenous infusions of blood-clotting factor as his ankle joint bleeds spontaneously. Before the age of five, the traumatic experience of being frequently injected was done in hospital by doctors or nurses. After he turned five, his parents learnt to administer the infusions at home. The ability to self-infuse means he can treat bleeds immediately at home instead of going to the hospital to seek help.

When K.C. was young, his parents opted for treatment only when he had a bleeding episode instead of regular infusions which will help prevent spontaneous bleeds. This was because the infusions were very expensive. One vial of factor can cost from \$130 to over \$1000, depending on the sub-type of haemophilia.

Although K.C. is thin, he requires a minimum of five vials to reduce bleeding. Dosage depends on the patient’s body weight – the greater the weight, the larger the amount of factor needed. His family thus spends about \$1,250 a month on the infusions, after subsidies from the government and the Haemophilia Society of Singapore.

Haemophilia is a life-long disease and the high cost of treatment is a major issue. Insurance companies have also rejected K.C.’s parents’ attempts to buy him medical insurance.

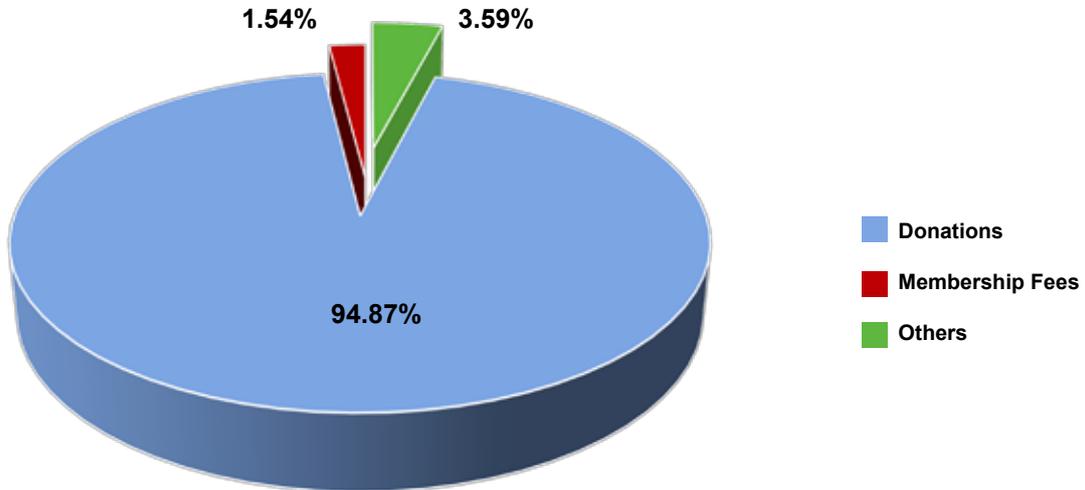
“Now that I am bigger, it is somewhat easier to cope with haemophilia as I am able to take care of myself and suffer fewer bleeding episodes,” says K.C. Currently he does not play any sports, but he knows how to swim and he enjoys playing the piano.

Thankfully, haemophilia has not affected his studies or social life. “I am like everyone else and I hang out with my friends.”

* Source: *Haemophilia Society of Singapore’s brochure*

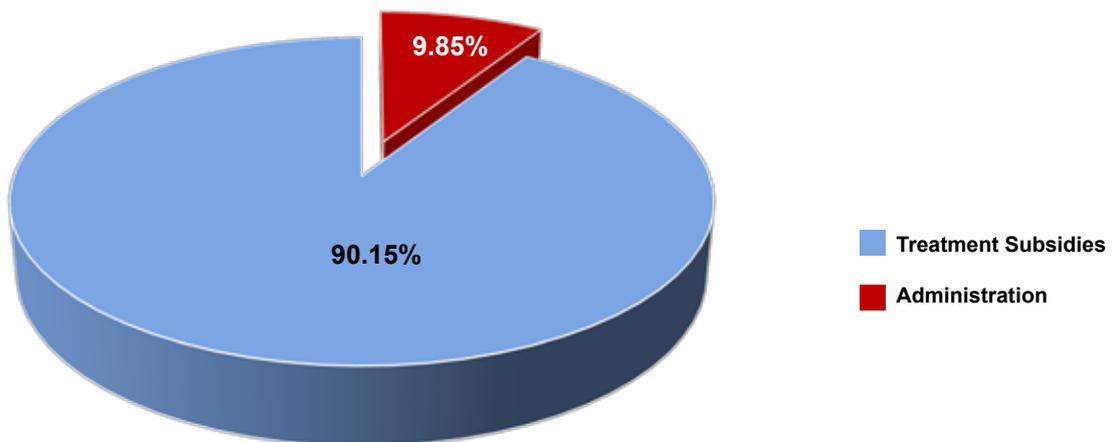
FINANCIAL INFORMATION SUMMARY

Breakdown of Total Income 2013/14



Donations	\$213,439	94.87%
Membership Fees	\$3,460	1.54%
Others	\$8,076	3.59%

Breakdown of Total Expenses 2013/14



Treatment Subsidies	\$153,556	90.15%
Administration	\$16,783	9.85%

THE HAEMOPHILIA SOCIETY OF SINGAPORE
(Charity Registration No: 00374 & Society No: S85SS0047A)

STATEMENT OF FINANCIAL POSITION AS AT 30 JUNE 2014

	<u>Note</u>	<u>2014</u> \$	<u>2013</u> \$
ASSETS			
Non-current assets			
Property, plant and equipment	4	259	379
Current assets			
Cash and bank balances	5	1,171,212	1,138,636
Grant receivable	7	21,400	-
Other receivables	6	1,600	1,960
		<u>1,194,212</u>	<u>1,140,596</u>
Total assets		<u>1,194,471</u>	<u>1,140,975</u>
EQUITY AND LIABILITIES			
Current liabilities			
Other payables	8	4,800	6,360
Deferred interest income		1,800	-
		<u>6,600</u>	<u>6,360</u>
Funds			
Unrestricted			
- Haemophilia welfare fund	9	50,265	52,065
- Dulcie Wilson scholarship fund	10	20,000	20,000
- Accumulated fund		1,117,606	1,062,550
		<u>1,187,871</u>	<u>1,134,615</u>
Total liabilities and funds		<u>1,194,471</u>	<u>1,140,975</u>

See accompanying notes to the financial statements

In our opinion, the financial statements are properly drawn up in accordance with the Charities Act, Cap 37, Recommended Accounting Practice 6, Societies Act, Cap 311 and Singapore Financial Reporting Standards and so as to give a true and fair view of the state of affairs of the society as at 30 June 2014 and the results, changes in funds and cash flows of the society for the financial year ended on that date.

Report on Other Legal and Regulatory Requirements

In our opinion, the accounting and other records required by the Act to be kept by the society have been properly kept in accordance with the provisions of the Act.

TKNP INTERNATIONAL
Public Accountants and
Chartered Accountants Singapore



STATEMENT OF FINANCIAL ACTIVITIES FOR THE FINANCIAL YEAR ENDED 30 JUNE 2014

Note	2014				2013			
	Unrestricted Funds	Restricted Funds	Endowment Funds	Total	Unrestricted Funds	Restricted Funds	Endowment Funds	Total
	\$	\$	\$	\$	\$	\$	\$	\$
Incoming resources								
<i>Incoming resources from generated funds</i>								
- Voluntary income	195,499	-	-	195,499	78,916	-	-	78,916
- Investment income	8,496	-	-	8,496	9,769	-	-	9,769
- Grant income	21,400	-	-	21,400	-	-	-	-
Total incoming resources	225,395	-	-	225,395	88,685	-	-	88,685
Resources expended								
<i>Cost of generating voluntary funds</i>								
- Depreciation	120	-	-	120	120	-	-	120
- Charity event	2,025	-	-	2,025	-	-	-	-
- Entertainment and refreshment	1,754	-	-	1,754	1,055	-	-	1,055
- General expenses	135	-	-	135	-	-	-	-
- IT related costs	1,613	-	-	1,613	754	-	-	754
- Miscellaneous	742	-	-	742	232	-	-	232
- Postage and stationery	3,934	-	-	3,934	1,524	-	-	1,524
	10,323	-	-	10,323	3,685	-	-	3,685
<i>Cost of charitable activities</i>								
- Medical fee subsidies to members	450	-	-	450	-	-	-	-
- Membership fees - NCSS	54	-	-	54	54	-	-	54
- Membership fees - WFH	557	-	-	557	-	-	-	-
- Sub-total carried forward	1,061	-	-	1,061	54	-	-	54

STATEMENT OF FINANCIAL ACTIVITIES FOR THE FINANCIAL YEAR ENDED 30 JUNE 2014

Note	2014				2013			
	Unrestricted Funds	Restricted Funds	Endowment Funds	Total	Unrestricted Funds	Restricted Funds	Endowment Funds	Total
	\$	\$	\$	\$	\$	\$	\$	\$
- Sub-total brought forward	1,061	-	-	1,061	54	-	-	54
- Subsidy of treatment – NUH Endowment Fund	77,320	-	-	77,320	89,041	-	-	89,041
- Subsidy of treatment – SGH Endowment Fund	32,700	-	-	32,700	40,860	-	-	40,860
- Subsidy of treatment – KKH Endowment Fund	39,150	-	-	39,150	27,372	-	-	27,372
- Welfare – outing for members	825	-	-	825	-	-	-	-
- Welfare – merit award	2,500	-	-	2,500	2,600	-	-	2,600
	153,556	-	-	153,556	159,927	-	-	159,927
<i>Governance costs</i>								
- Accountancy fees	3,200	-	-	3,200	3,100	-	-	3,100
- Auditors' remuneration	3,000	-	-	3,000	3,000	-	-	3,000
	6,200	-	-	6,200	6,100	-	-	6,100
<i>Finance cost</i>	260	-	-	260	465	-	-	465
Total resources expended	170,339	-	-	170,339	170,177	-	-	170,177
Net surplus/(deficit) resources	55,056	-	-	55,056	(81,492)	-	-	(81,492)

LIST OF DONORS

INDIVIDUALS

Ang Edwin	Khoo Chwee Tin	Seah Yuri
Ang Han Chunn, Dannyn	Kinman, John	See Beng Teck (Dr)
Ang Peng Chye (Dr)	Koh Barbara	See Li Ling
Ang Shu Qiong	Koh Zhen Hao	Seng Ah Ngoh, Nerine
Au Ying Lai, Lydia	Kong Wai Ping, Irene	Seng Boon Kheng (Dr)
Chan Chew Hua	Kum Kelly	Sia Yoke Leng
Chai Siow Peng	Kuo Yew Ting	Sim Wee Li
Chan Siew Hoon	Lam Ching Mei, Joyce (Dr)	Soeparto
Chee Chun Woei	Lam Yeng Chin	Soh Siew Hong
Chee Jing Jye (Dr)	Le Meur, Carole	Song Darelle
Chee Liew Chin	Lee Hwee Chin, Erin	Tan Hooi Hwa (Dr)
Chen Wei Ching	Lee Michelle	Tan Huang Tai, Peter
Chew Hwee Koon, Ivy	Lee Ren Sheng	Tan Isobel
Chin Wai Huoy, Claire	Liaw Jimmy	Tan Keshia
Choe Peng Sum	Lim Ah Lay, Judy	Tan Kim Hock
Choh Siew Choon	Lim Cheng Bee	Tan Lai Ean
Chong Gabriel	Lim Chiew Har	Tan Lai See
Chong Siew Peng	Lim Chin Yiong, Jesper	Tan Loon Ying
Chong Yee Feng	Lim Lian Arn (Dr)	Tan Siew Tuan
Chu Kim Mui	Loh Sow Mui, Karin	Tan Suat Cheng
Chua Chung Peow	Low Kok Chun	Tan Wai Peng
Chua Han Yong	Low Li Ying	Tang Wee Loke
Chua Harry	Low Wai Yee, Yvonne	Tensingh, Edwin
Chua Meng Pin	MayKyaw Soe Nyunt	Teo Soo Ling, Catherine
Dharmadi Betty	Ng Geok Hoon, Agnes	Tham Chuey Ping
Foo Cui Yun	Ng Jolene	Tien Sim Leng (Dr)
Foo Siang-Kee	Ng Leong Chin	Toh Wei Lin
Fu Chun Yan	Ng Mary	Tong Moo Kiat
Gan Rosie	Ng Peik Lan	Vaz, Gerard
Gehle, Helmut	Ng Shuh Fang	Wagner, Juergen
Ghui Zhong Jun, Shannon	Ng Teck Hiang	Wee Kim Hin
Goh Han Meng	Ng Wei Ren, Clement	Wee Shen Wen, Clinton
Goh Koon Tho	Nur Sakinah bte Mahadi	Wee Shiang Ning
Goh Poh Ngee	Ong Eng Hwa, Christopher	Wee Soon Bing
Goh Siew Hong	Ong Lay Har	Whang Shang Ying
Goh Siong Hoe	Ong Yeow Chon	Whang Tar Liang
Hall, Carolyn	Pan Cheng Lui	Wong Chen Liong
Heah Siew Min	Pan Yi Wen, Andy	Wong Chi Lun, Allan
Heng Puay Fung	Pang Heng Mun	Wong George
Ho Kah Chung	Poh Beow Kiong	Wong Kee Ming
Ho Kwong Soon	Pon Yuet Peng	Xiao Zhiqiang, Zed
Ho May Sian	Quek Hwee Choo	Yap Boon Kim, Pamela
Hong Siew Chen	Quek Sim Pin	Yip Lai Lai, Lily
Hu Ian Jin	Rajah, Chelva	Yong Kuek Siong (Dr)
Kaul, Shankar	Raull, Mary	Young, Richard Fe

ORGANISATIONS

AllAlloy Dynaweld Pte Ltd	Loyang Tua Pek Kong Temple
Baxter Healthcare (Asia) Pte Ltd	Qeren Communications Pte Ltd
Char Yong (DABU) Fdn Pte Ltd	Royal Thai Embassy
DCA Architects Pte Ltd	Sartorius Stedim Pte Ltd
Eshcol Pharmaceutical Group	Singapore Island Country Club
Hillgrove Secondary School	Singapore Press Holdings Ltd
Lee Foundation	

The Society's Song

A RAY OF HOPE

Everyday as the world unfolds
We can see we are not alone
Hand in hand, everyone we stand
Firm with no regrets

Though at times, we say life is unfair
At times we just can't go on
Then again, just a few steps away
We witness a ray of hope

So be strong, let us carry on
To live a life worth living for
Travelling on with our hearts as one
Never letting go

So at times, we say life is unfair
Remember our song, our prayer
Let it be our pillar of strength
For many more years to come

Mmm.....(fade to end)

This song was composed by Dayvid Foo,
a member of the Haemophilia Society of Singapore.
The song can be accessed from the Society's website.

Annual Report Editorial Committee

Dr Tan Hooi Hwa
Dr Gan Kim Loon
Wee Ai Choo
See Ek May