

1. Company details

Name of entity

CYCLOPHARM LIMITED

ABN or equivalent company reference	Financial year ended ('current period')	Financial year ended ('previous period')
74 116 931 250	31 December 2010	31 December 2009

2. Results for announcement to the market

2.1 Revenues from ordinary activities	down	(14.7%)	to	9,464,605
2.2 Profit from ordinary activities after tax attributable to members	down	(78.0%)	to	450,106
2.3 Net profit for the period attributable to members	down	(78.0%)	to	450,106
2.4 Dividends	Amount per security		Franked amount per security	
Final dividend proposed	Not applicable		Not applicable	
Interim dividend	Not applicable		Not applicable	
2.5 Record date for determining entitlements for the final dividend	<div style="border: 1px solid black; display: inline-block; padding: 5px 20px;">Not applicable</div>			
2.6 Brief explanation of any of the figures in 2.1 to 2.4 above necessary to enable the figures to be understood.				
Refer Attachment 1.				

3. Statement of financial performance

Refer Attachment 1.

4. Statement of financial position

Refer Attachment 1.

5. Statement of cash flows

Refer Attachment 1.

6. Dividends

Not applicable

7. Dividend reinvestment plans

Not applicable

8. Statement of retained earnings

Refer Attachment 1.

9. Net tangible assets

Refer Attachment 1.

10. Entities over which control has been gained or lost during the period

Control over entities

Name of entity (or group of entities)

Refer Attachment 1.

Loss of control over entities

Name of entity (or group of entities)

Refer Attachment 1.

11. Details of associates and joint venture entities

Refer Attachment 1.

12. Significant Information

Refer Attachment 1.

13. Foreign Entities

Refer Attachment 1.

14. Commentary on results for the period

Refer Attachment 1.

15. A statement as to whether the report is based on accounts which have been audited or subject to review, are in the

The accounts are in the process of being audited.

16. If the accounts have not yet been audited or subject to review and are likely to be subject to dispute or qualification, details are described below

The accounts are unlikely to be subject to dispute or qualification.

17. If the accounts have been audited or subject to review and are subject to dispute or qualification, details are described below

Not applicable

Contact details:

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Company Secretary
Cyclopharm Limited

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Appendix 4E
Preliminary Final Report
For the year ended 31 December 2010

Cyclopharm Limited and its Controlled Entities
ABN 74 116 931 250

cyclopharm

Managing Director's Report

Continued



Appendix 4E Commentary Full Year Results of Cyclopharm Limited and its Controlled Entities ("Company") For the 12 months ended 31 December 2010

Features

I am pleased to provide you with the overview of Cyclopharm Limited's ("Cyclopharm") performance for 2010. Since listing in 2007, this past year will represent a significant year for Cyclopharm with the commercialisation of key projects. The delivery of these projects have not only diversified your company but also opened a path for further expansion and profitability.

In the Company's listing Prospectus the Director's outlined a strategy to develop a new Molecular Imaging division to manufacture and sell radiopharmaceuticals including F18 Flurodeoxyglucose (FDG) for use in Positron Emission Tomography (PET) scanning. During the year, we commissioned our first PET facility at Macquarie University Hospital (MUH) and received a Good Manufacturing Process (GMP) license approval from the Australian Therapeutic Goods Administration (TGA).

The Molecular Imaging business contributed revenues of \$21,842 (2009: \$0) and a Loss after tax of \$864,385 (2009: \$250,755). During the year in review, property, plant and equipment increased to \$10,059,528 (2009: \$5,052,951) due primarily to the development of our PET nuclear pharmacy at MUH. The Director's consider the commercialisation of the Molecular Imaging division as a key step in increasing shareholder value.

Macquarie Medical Imaging (MMI) purchased our first commercial doses of FDG. MMI, Cyclopharm's medical imaging joint venture with Alfred Health Solutions provides state of the art imaging facilities including 3T MRI, CT, X-ray, Ultrasound and PET scanning for the new hospital located on the campus of Macquarie University and neighbouring suburbs. During the year, Cyclopharm made an investment of \$500,000 into MMI (2009: \$0). The share of the loss in associate was \$311,372 (2009: \$0).

The combined sales of the Company's key products TechnegasPlus generators (Generators) and Patient Administration Sets (PAS), decreased from \$11.09m to \$9.46m or 15%. Techengas Generator revenue was consistent with the prior year while both the sale of Patient Administration Sets (PAS) boxes and revenues were lower.

As mentioned in the 2009 Annual Report and 2010 Half Year Report, PAS sales were dampened by the global molybdenum shortage. Molybdenum is used to make Technetium⁹⁹ which is an essential component of Technegas. In addition unfavourable foreign currency movements impacted foreign sales.

For the reasons mentioned above, the Company's profit after tax of \$450,106 (2009: \$2,044,490) was negatively impacted. The table below outlines the current year to prior year results whereby non-recurring revenues and costs relating to a case against Clinquest Inc, the share of the loss in associate and Molecular imaging division are adjusted. A normalised comparison follows:

Full Year ending 31 December		2009	2010
Net Profit Before Tax	\$'000	2,568	808
Clinquest settlement net gain	\$'000	(524)	-
MMI share of loss of associate	\$'000	-	311
Molecular Imaging Division loss	\$'000	-	864
Adjusted Net Profit Before Tax		2,044	1,983

Managing Director's Report

Continued



Operating review

Technegas

Technegas is a lung imaging device used primarily to diagnose the presence of blood clots in the lungs known as Pulmonary Emboli (PE). For the last 20 years, over 2,400,000 patients have benefited from the Technegas system. The company faced similar challenges as in prior years with continued competition from Computed Tomography Pulmonary Angiogram ("CTPA") and the global molybdenum shortage dampened demand for all nuclear medicine products. In spite of these hurdles Technegas has demonstrated its relevance and has once again generated sales and cash flows.

Revenue Composition

Overall sales revenue of \$9.46m from the Company's key products, Generators and PAS were lower than the preceding year (2009: \$11.09m). PAS or consumable revenue declined 19% to \$7.36 million (162,800 units) for the current period compared to that of the previous year (2009: \$9.13 million or 163,250 units). Sales margins were significantly lower than the prior year due to foreign exchange movements in the Euro to Australian dollar and a change in sales by region.

We recorded 54 Technegas Plus Generator sales in 2010, which is consistent with the prior year (2009: 57). Sales into Asia were strong with 12 generators being sold into China during the year. Technegas generator and other sales revenues of \$2.08m were consistent with the prior year (2009: \$1.96m). The strengthening of the Australian dollar against the Euro impacted revenues and gross margins as approximately 60% of sales are made to European countries.



New Drug Application to sell Technegas in the USA

As required by the FDA, the Company's next step is to undertake an additional Phase III clinical trial prior to approval of Technegas for sale in the USA. After a series of proposals and meetings throughout 2010, the FDA in February 2011 approved the Company's Special Protocol Assessment (SPA).

A SPA is a mechanism through which the FDA and sponsor reach agreement on the design, size, clinical endpoints, and data analysis of a clinical trial that is intended to support an efficacy claim in a New Drug Application (NDA) for regulatory approval. The SPA ensures that the agreed clinical trial design meets the FDA's expectations for a pivotal study.

Over half of the world's nuclear medicine departments are located in the United States. Although, not all nuclear medicine departments perform lung imaging, the significance of this market for Technegas is undeniable. It has been a strategy of the Company to obtain approval to sell Technegas for over 10 years.

In parallel with the clinical trial development, we have also engaged a number of the leading United States medical centres to participate in the upcoming trial. Technegas is the globally recognised ventilation imaging agent of choice in determining pulmonary embolism from both a safety and efficacy standpoint. While we will only require 10 to 15 centres for our clinical trial, we have been approached by scores more wanting to be first to have access to our technology in the United States.

Managing Director's Report Continued

New Drug Application to sell Technegas in the USA (continued)

Potential clinical trial sites are now submitting budgets for the trial. Provided clinical site negotiations go well, patient recruitment is anticipated to start mid 2011. It is expected that the clinical trial will take a year to complete followed by a New Drug Application submission to the FDA. Based on this timeline, it is anticipated that Technegas will be available for sale in the United States in 2014. As the USA represents such a major growth opportunity, your Directors will continue to sensibly pursue entry into this market.

Regional Review

Europe

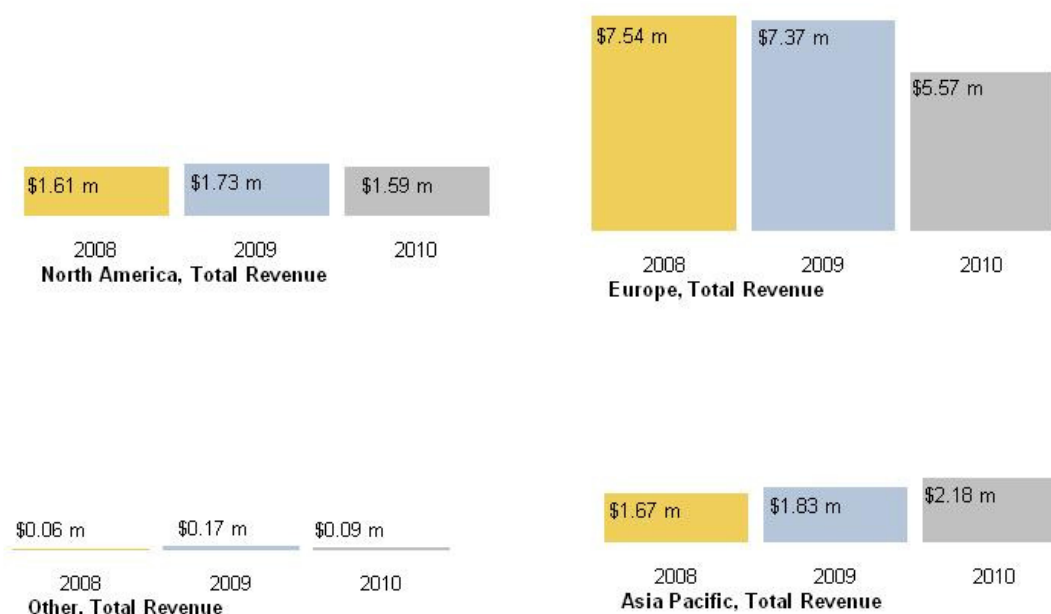
Approximately 60% of sales revenue is derived in Europe compared with 68% in 2009. Sales revenue decreased 24% on the same time last year despite higher Generator sales. 30 Generators were sold in 2010 compared with 24 in the prior year. Revenues from PAS sales were down 30% lower on 2009 due changes in the sales mix (fewer sales were made to countries with high gross margins). Sales revenue was also impacted by an approximate 15% movement in the Australian dollar relative to the Euro.

North America

In the 2009 Annual Report we advised shareholders that North America revenues were expected to grow by 4% however revenues from the sale of Generators and PAS decreased 9%, on the prior year. Only 4 generators were sold in 2010 (2009: 12) while a 7th year of consecutive growth in PAS unit sales were recorded with 658 PAS boxes sold (2009: 585 PAS boxes). Total revenue of \$1.59m has meant Canada is now our second largest market behind Europe. The directors have been overjoyed with the success of Technegas in Canada. Now that the FDA has approved our SPA we are well positioned to leverage from our Canadian team once approval to sell Technegas is obtained.

Asia Pacific

Revenues in the Asia Pacific region were 19% higher than 2009. In Australia, where Technegas enjoys a very high market share total revenues grew by 9%. In Asia, sales almost doubled. Sales to Hong Kong and China drove the increase in revenues of 101% over the prior year. This trend is not expected to continue at such a rapid rate however sales executives will focus on expanding the footprint of Technegas in leading hospitals throughout China's mainland and surrounding territories. The new regulatory approval in Japan was not received during 2010 and we expect sales growth once approval is received.



Managing Director's Report

Continued



Molecular Imaging

In the Company's listing Prospectus the strategy to establish a new revenue stream for the Company, in the form of the Molecular Imaging division was outlined. The premise for the new division was the growth in the incidence of Australians affected by cancer (1 in 3 males and 1 in 4 females before the age of 75) and the impact PET (Positron Emission Tomography) imaging was having in Europe and the United States on cancer sufferers.

PET is clinically proven to better identify the location and extent of certain active cancer cells in the body and to assist physicians to refine the course of resection or therapy. The Directors maintain their vision to establish PET Facilities in major Australian capital cities which will supply PET radiopharmaceuticals to hospitals for use in PET scans.

In July 2010, MUH's 183 bed facility was opened. The facility will support 12 operating theatres and is designed to be a clinical centre of excellence in oncology and neurology. It is Australia's first privately owned, campus-based university hospital and will combine academic medicine with group practice. As Australia's newest university hospital, MUH will support the advancement of future surgeons and leaders of the medical profession.

In December 2010 Cyclopharm announced to its Shareholders that the wholly owned subsidiary Cyclopet Pty Limited (Cyclopet) received Good Manufacturing Process (GMP) license approval from the Australian Therapeutic Goods Administration (TGA) for the cyclotron production and research facility located at Macquarie University Hospital (MUH), our first Molecular Imaging facility.

In December 2010, Cyclopharm advised Shareholders that the first commercial doses of the radiopharmaceutical compound F18 Flurodeoxyglucose (FDG) for use in Positron Emission Tomography (PET) scanning were sold to MMI. PET is the fastest growing nuclear medicine diagnostic imaging modality in the world. PET is clinically proven to better identify the location and extent of certain active cancer cells in the body as well as many neurodegenerative diseases.

Macquarie Medical Imaging

Macquarie Medical Imaging commenced its operations in May 2010 just prior to the grand opening of Macquarie University in July 2010 and purchased the first commercial doses of FDG. The first patient injected with our product confirmed the presence and location of Squamous Cell Carcinoma (SCC). SCC is a form of cancer of the carcinoma type that may occur in many different organs, including the skin, lips, mouth, esophagus, urinary bladder, prostate, lungs, vagina, and cervix. 115,000 Australians will be diagnosed with cancer this year. This figure represents a 10% increase since 2006. Cyclopet's products and research will play a major role in early diagnosis and therapeutic management of these cancer patients.

MMI, Cyclopharm's medical imaging joint venture with Alfred Health Solutions provides access to state of the art imaging facilities including 3T MRI, CT, X-ray, Ultrasound and PET scanning for the new hospital and neighbouring suburbs.

PET radiopharmaceuticals will be supplied on-site to Macquarie Medical Imaging and other hospitals predominately located in New South Wales. Other revenue streams generated from the PET radiopharmaceutical production facility include opportunities for Phase III clinical trials and pharmaceutical drug development.

Your directors are pleased with the quality of the MMI facility and hold the strong belief in the synergies that the joint venture and the co-located PET facility will provide such as the provision of a fully aligned and integrated diagnostic, therapeutic and research platform.

Managing Director's Report

Continued



OUTLOOK

In 2011, your Directors expect growth in Technegas revenues to result from targeted marketing and new anticipated regulatory approvals campaigns in Japan, Russia and South America. We forecast a change in the mix of Technegas products. We anticipate more generators (lower margins) relative to PAS box sales and therefore lower profit margins.

Your directors expect revenues from the PET Facility at the MUH to ramp up throughout the year and for the division to be profitable by year's end. We expect operating costs, depreciation and finance costs to be higher since the Molecular Imaging division will be operational for the full 12 months.

The Director's maintain their view that the PET facility is a major investment that will yield significant long term returns for the Company but recognise that the interim working capital shortfall will require funding from the Technegas business.

We expect to make a further investment into MMI of circa \$0.30m during the year to part-fund the working capital requirements of the business during its preliminary stages. Our estimates indicate the venture will generate a substantial return on investment in subsequent years.

James McBrayer
Managing Director and CEO

Statement of Comprehensive Income

for the year ended 31 December 2010



	Notes	Consolidated	
		2010 \$	2009 \$
CONTINUING OPERATIONS			
Sales revenue	4	9,464,605	11,098,579
Finance revenue		68,972	93,186
Other revenue	4	-	2,403,090
Total revenue		9,533,577	13,594,855
Cost of materials and manufacturing	4a	(2,308,992)	(2,546,484)
Employee benefits expense	4e	(2,957,231)	(3,537,984)
Advertising and promotion expense		(238,646)	(271,165)
Depreciation and amortisation expense	4c	(350,546)	(450,229)
Freight and duty expense		(342,666)	(384,682)
Research and development expense	4d	(16,847)	(39,728)
Administration expense	4f	(1,663,503)	(2,875,245)
Other expenses	4g	(337,067)	(784,334)
Share of loss of an associate	11	(311,372)	-
Profit / (loss) before tax and finance costs		1,006,707	2,705,004
Finance costs	4b	(198,299)	(137,503)
Profit / (loss) before income tax		808,408	2,567,501
Income tax expense	5	(358,302)	(523,011)
Net profit for the year		450,106	2,044,490
Other comprehensive income after income tax			
Exchange differences on translating foreign controlled entities		(708,770)	(1,363,434)
Total comprehensive (loss) / income for the year		(258,664)	681,056
Earnings per share (cents per share)	6	cents	cents
-basic earnings per share for continuing operations		0.26	1.20
-basic earnings per share		0.26	1.20
-diluted earnings per share		0.26	1.20

The Statement of Comprehensive Income is to be read in conjunction with the notes to the financial statements.

Statement of Financial Position

as at 31 December 2010

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Nuclear Medicine



Consolidated			
	Notes	2010 \$	2009 \$
Assets			
Current Assets			
Cash and cash equivalents	7	1,541,644	4,612,205
Trade and other receivables	8	4,295,487	4,281,369
Inventories	9	2,546,867	3,242,992
Other assets		25,490	33,227
Total Current Assets		8,409,488	12,169,793
Non-current Assets			
Inventories	9	702,000	-
Trade and other receivables	8	161,500	-
Property, plant and equipment	10	10,059,528	5,052,951
Investments in associate	11	188,231	-
Intangible assets	12	2,621,152	2,421,667
Total Non-current Assets		13,732,411	7,474,618
Total Assets		22,141,899	19,644,411
Liabilities			
Current Liabilities			
Trade and other payables	13	1,082,492	1,900,153
Interest bearing loans and borrowings	14	6,078,300	1,757,350
Provisions	15	499,283	498,283
Tax liabilities	5	318,850	174,039
Total Current Liabilities		7,978,925	4,329,825
Non-current Liabilities			
Interest bearing loans and borrowings	14	24,453	975,900
Provisions	15	27,710	42,741
Deferred tax liabilities	5	344,657	367,304
Total Non-current Liabilities		396,820	1,385,945
Total Liabilities		8,375,745	5,715,770
Net Assets		13,766,154	13,928,641
Equity			
Contributed equity	16	11,088,908	11,088,908
Employee equity benefits reserve	23	314,149	217,972
Foreign currency translation reserve		(1,543,224)	(834,454)
Retained Profits / (Accumulated losses)		3,906,321	3,456,215
Total Equity		13,766,154	13,928,641

The Statement of Financial Position is to be read in conjunction with the notes to the financial statements.

Statement of Cash Flows

for the year ended 31 December 2010

		Consolidated	
		2010	2009
		\$	\$
Notes			
Operating activities			
	Receipts from customers	9,458,224	14,569,018
	Payments to suppliers and employees	(9,183,719)	(10,641,452)
	Interest received	68,972	93,186
	Borrowing costs paid	(198,299)	(137,503)
	Income tax paid	(236,138)	(192,216)
	Net cash flows from / (used) operating activities	(90,960)	3,691,033
Investing activities			
	Investment in associate	(500,000)	-
	Purchase of property, plant and equipment	(5,470,211)	(2,732,029)
	Payments for deferred expenditure	(247,897)	(364,836)
	Net cash flows used in investing activities	(6,218,108)	(3,096,865)
Financing activities			
	Proceeds from borrowings	3,743,603	-
	Repayment of borrowings	(374,100)	-
	Net cash flows from financing activities	3,369,503	-
	Net increase / (decrease) in cash and cash equivalents	(2,939,565)	594,168
Cash and cash equivalents			
	- at beginning of the period	4,612,205	4,206,271
	- net foreign exchange differences from translation of cash and cash equivalents	(130,996)	(188,234)
	- at end of the period	1,541,644	4,612,205

The Statement of Cash Flows is to be read in conjunction with the notes to the financial statements.

Statement of Changes in Equity

for the year ended 31 December 2010



	Share capital	Other Contributed Equity	Total Contributed Equity	Accumulated Profits	Foreign Currency Translation Reserve	Employee Equity Benefits Reserve	Attributable to Equity Holders of the Parent	Total
	\$	\$	\$	\$	\$	\$	\$	\$
CONSOLIDATED								
Balance at								
1 January 2009	16,422,066	(5,295,657)	11,126,409	1,411,725	528,980	143,689	13,210,803	13,210,803
Cost of share based payments	-	-	-	-	-	74,283	74,283	74,283
Other comprehensive income	-	-	-	-	(1,363,434)	-	(1,363,434)	(1,363,434)
Total income (expense) for the year recognised directly in equity	-	-	-	-	(1,363,434)	74,283	(1,289,151)	(1,289,151)
Profit for the year	-	-	-	2,044,490	-	-	2,044,490	2,044,490
Total income (expense) for the year	-	-	-	2,044,490	(1,363,434)	74,283	755,339	755,339
Other	-	(37,501)	(37,501)	-	-	-	(37,501)	(37,501)
Balance at								
31 December 2009	16,422,066	(5,333,158)	11,088,908	3,456,215	(834,454)	217,972	13,928,641	13,928,641
Balance at								
1 January 2010	16,422,066	(5,333,158)	11,088,908	3,456,215	(834,454)	217,972	13,928,641	13,928,641
Cost of share based payments	-	-	-	-	-	96,177	96,177	96,177
Other comprehensive income	-	-	-	-	(708,770)	-	(708,770)	(708,770)
Total income (expense) for the year recognised directly in equity	-	-	-	-	(708,770)	96,177	(612,593)	(612,593)
Profit for the year	-	-	-	450,106	-	-	450,106	450,106
Total income (expense) for the year	-	-	-	450,106	(708,770)	96,177	(162,487)	(162,487)
Balance at								
31 December 2010	16,422,066	(5,333,158)	11,088,908	3,906,321	(1,543,224)	314,149	13,766,154	13,766,154

The Statement of Changes in Equity is to be read in conjunction with the notes to the financial statements.

Notes

for the year ended 31 December 2010

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1. CORPORATE INFORMATION

Cyclopharm is a Company limited by shares incorporated and domiciled in Australia. The shares are publicly traded on the Australian Securities Exchange ("ASX").

During the year the principal continuing activities of the consolidated entity consisted of the manufacture and sale of medical equipment and radiopharmaceuticals, including associated research and development.

2. SUMMARY OF SIGNIFICANT ACCOUNTING POLICIES

a) Basis of Preparation

The financial report is a general-purpose financial report, which has been prepared in accordance with the requirements of the Corporations Act 2001 and Australian Accounting Standards. The financial report has been prepared on a historical cost basis.

The financial report is presented in Australian dollars.

b) Statement of compliance

The financial report complies with Australian Accounting Standards, which include Australian equivalents to International Financial Reporting Standards ('AIFRS'). Compliance with AIFRS ensures that the financial report, comprising the financial statements and notes thereto, complies with International Financial Reporting Standards ('IFRS').

The following standards and amendments were available for early adoption but have not been applied by the consolidated entity in these financial statements:

New accounting standards and interpretations not yet adopted

There are no changes to accounting policies applicable for the financial year ended 31 December 2010 for the Company and the consolidated entity.

The following Australian Accounting Standards have been issued or amended and are applicable to the parent and consolidated entity but are not yet effective. They have not been adopted in preparation of the financial statements at reporting date.

The AASB has issued new, revised and amended standards and interpretations that have mandatory application dates for future reporting periods. The Group has decided against early adoption of these standards. A discussion of those future requirements and their impact on the Group follows:

AASB 9: Financial Instruments and AASB 2009–11: Amendments to Australian Accounting Standards arising from AASB 9 [AASB 1, 3, 4, 5, 7, 101, 102, 108, 112, 118, 121, 127, 128, 131, 132, 136, 139, 1023 & 1038 and Interpretations 10 & 12] (applicable for annual reporting periods commencing on or after 1 January 2013).

These Standards are applicable retrospectively and amend the classification and measurement of financial assets. The Group has not yet determined potential impact on the financial statements.

The changes made to accounting requirements include:

- simplifying the classifications of financial assets into those carried at amortised cost and those carried at fair value;
- simplifying the requirements for embedded derivatives;
- removing the tainting rules associated with held-to-maturity assets;
- allowing an irrevocable election on initial recognition to present gains and losses on investments in equity instruments that are not held for trading in other comprehensive income. Dividends in respect of these investments that are a return on investment can be recognised in profit or loss and there is no impairment or recycling on disposal of the instrument; and

- requiring financial assets to be reclassified where there is a change in an entity's business model as they are initially classified based on: (a) the objective of the entity's business model for managing the financial assets; and (b) the characteristics of the contractual cash flows.

AASB 1053: Application of Tiers of Australian Accounting Standards and AASB 2010–2: Amendments to Australian Accounting Standards arising from Reduced Disclosure Requirements [AASB 1, 2, 3, 5, 7, 8, 101, 102, 107, 108, 110, 111, 112, 116, 117, 119, 121, 123, 124, 127, 128, 131, 133, 134, 136, 137, 138, 140, 141, 1050 & 1052 and Interpretations 2, 4, 5, 15, 17, 127, 129 & 1052] (applicable for annual reporting periods commencing on or after 1 July 2013).

AASB 1053 establishes a revised differential financial reporting framework consisting of two tiers of financial reporting requirements for those entities preparing general purpose financial statements:

Tier 1: Australian Accounting Standards; and

Tier 2: Australian Accounting Standards — Reduced Disclosure Requirements.

This Standard deems the Group to be a Tier 1 entity and hence there is no accounting impact to be considered going forward.

AASB 2009–10: Amendments to Australian Accounting Standards — Classification of Rights Issues [AASB 132] (applicable for annual reporting periods commencing on or after 1 February 2010).

This Standard clarifies that rights, options or warrants to acquire a fixed number of an entity's own equity instruments for a fixed amount in any currency are equity instruments if the entity offers the rights, options or warrants pro rata to all existing owners of the same class of its own non-derivative equity instruments. The amendments are not expected to have significant impact on the Group.

AASB 2009–12: Amendments to Australian Accounting Standards [AASBs 5, 8, 108, 110, 112, 119, 133, 137, 139, 1023 & 1031 and Interpretations 2, 4, 16, 1039 & 1052] (applicable for annual reporting periods commencing on or after 1 January 2011).

This Standard makes a number of editorial amendments to a range of Australian Accounting Standards and Interpretations, including amendments to reflect changes made to the text of IFRSs by the IASB. The Standard also amends AASB 8 to require entities to exercise judgment in assessing whether a government and entities known to be under the control of that government are considered a single customer for the purposes of certain operating segment disclosures.

The amendments are not expected to impact the Group.

AASB 2009–13: Amendments to Australian Accounting Standards arising from Interpretation 19 [AASB 1] (applicable for annual reporting periods commencing on or after 1 July 2010).

This Standard makes amendments to AASB 1 arising from the issue of Interpretation 19. The amendments allow a first-time adopter to apply the transitional provisions in Interpretation 19.

This Standard is not expected to impact the Group.

AASB 2009–14: Amendments to Australian Interpretation — Prepayments of a Minimum Funding Requirement [AASB Interpretation 14] (applicable for annual reporting periods commencing on or after 1 January 2011).

This Standard amends Interpretation 14 to address unintended consequences that can arise from the previous accounting requirements when an entity prepays future contributions into a defined benefit pension plan. This Standard is not expected to impact the Group.

AASB 2010–3: Amendments to Australian Accounting Standards arising from the Annual Improvements Project [AASB 3, AASB 7, AASB 121, AASB 128, AASB 131, AASB 132 & AASB 139] (applicable for annual reporting periods commencing on or after 1 July 2010).

- requiring that recognition and/or adjustment of contingent consideration for acquisitions preceding 1 July 2009 be recognised against the cost of acquisition
- clarifying the accounting for replacement share-based payments awarded to the acquiree's employees as part of the cost of the combination service, or in the case of non-replaced and unvested share-based payments of the acquiree that do not form part of the exchange, an allocation to both the cost of acquisition and post-combination services on the basis of a market-based measure; and
- making sundry transitional amendments to various Standards

This Standard is not expected to impact the Group.

AASB 2010–4: Further Amendments to Australian Accounting Standards arising from the Annual Improvements Project [AASB 1, AASB 7, AASB 101 & AASB 134 and Interpretation 13] (applicable for annual reporting periods commencing on or after 1 January 2011).

This Standard details numerous non-urgent but necessary changes to Accounting Standards arising from the IASB's annual improvements project. Key changes include:

- clarifying the application of AASB 108 prior to an entity's first Australian-Accounting-Standards financial statements;
- adding an explicit statement to AASB 7 that qualitative disclosures should be made in the context of the quantitative disclosures to better enable users to evaluate an entity's exposure to risks arising from financial instruments;
- amending AASB 101 to the effect that disaggregation of changes in each component of equity arising from transactions recognised in other comprehensive income is required to be presented, but is permitted to be presented in the statement of changes in equity or in the notes;
- adding a number of examples to the list of events or transactions that require disclosure under AASB 134; and
- making sundry editorial amendments to various Standards and Interpretations.

This Standard is not expected to impact the Group.

AASB Interpretation 19: Extinguishing Financial Liabilities with Equity Instruments (applicable for annual reporting periods commencing from 1 July 2010).

This Interpretation deals with how a debtor would account for the extinguishment of a liability through the issue of equity instruments. The Interpretation states that the issue of equity should be treated as the consideration paid to extinguish the liability, and the equity instruments issued should be recognised at their fair value unless fair value cannot be measured reliably, in which case they shall be measured at the fair value of the liability extinguished. The Interpretation deals with situations where either partial or full settlement of the liability has occurred. This Interpretation is not expected to impact the Group.

The Group does not anticipate early adoption of any of the above reporting requirements and does not expect these requirements to have any material effect on the Group's financial statements.

c) Basis of consolidation

The consolidated financial statements comprise the financial statements of Cyclopharm and its subsidiaries as at 31 December each year ('the Group').

Subsidiaries

Subsidiaries are consolidated from the date on which control is transferred to the Group and cease to be consolidated from the date on which control is transferred out of the Group. Where there is loss of control of a subsidiary, the consolidated financial statements include the results for the part of the reporting period during which Cyclopharm has control.

The financial statements of subsidiaries are prepared for the same reporting period as the parent Company, using consistent accounting policies. Adjustments are made to bring into line any dissimilar accounting policies that may exist.

Transactions eliminated on consolidation

All intercompany balances and transactions, including unrealised profits arising from intra-group transactions, have been eliminated in full. Unrealised losses are eliminated unless costs cannot be recovered.

The Directors have identified that the business combination, encompassing the restructure of the Cyclopharm Group that occurred in May 2006 constituted a reverse acquisition as defined under *AASB3 Business Combinations*. Accordingly the consolidated financial statements have been issued under the name of the new legal parent, Cyclopharm, but reflect a continuation of the financial statements of the economic entity that existed prior to the business combination/reorganisation.

For business combinations involving entities under common control, which are outside the scope of *AASB 3 Business Combinations*, the Company applies the purchase method of accounting by the legal parent.

d) Foreign currency translation

Functional and presentation currency

The functional currency of each of the group's entities is measured using the currency of the primary economic environment in which that entity operates. The consolidated financial statements are presented in Australian dollars (Aud \$) which is the parent entity's functional and presentation currency.

Transactions and balances

Transactions in foreign currencies are initially recorded in the functional currency at the exchange rates ruling at the date of the transaction. Foreign currency monetary items are translated at the year-end exchange rate. Non-monetary items that are measured in terms of historical cost continue to be carried at the exchange rate at the date of the transaction. Non-monetary items measured at fair value are reported at the exchange rate when the fair value was determined.

Exchange differences arising on the translation of monetary items are recognised in the Statement of Comprehensive Income, except where deferred in equity as a qualifying cash flow or net investment hedge. On disposal of a foreign entity the deferred cumulative amount in equity is recognised in the Statement of Comprehensive Income.

Group companies

The functional currency of the overseas subsidiaries Cyclomedica Ireland Limited, Cyclomedica Germany GmbH, Cyclomedica Europe Limited, is European Euro (Euro €) and Cyclomedica Canada Limited is Canadian dollars (Can \$).

The financial results and position of foreign operations whose functional currency is different from the group's presentation currency are translated as follows:

- Assets and liabilities are translated at year-end exchange rates prevailing at that reporting date.
- Income and expenses are translated at the weighted average exchange rates for the period.
- Retained profits are translated at the exchange rates prevailing at the date of the transaction.

Exchange differences arising on the translation of foreign operations are transferred directly to the group's foreign currency translation reserve in the Statement of Financial Position. These differences are recognised in the Statement of Comprehensive Income in the period in which the entity is disposed. Exchange differences arising on the translation of non-monetary items are recognised directly in equity to the extent that the gain or loss is directly recognised in equity, otherwise the exchange difference is recognised in the Statement of Comprehensive Income.

e) Income tax

Income tax on the profit and loss for the year comprises current and deferred tax. Income tax is recognised in the Statement of Comprehensive Income, except to the extent that it relates to items recognised directly to equity, in which case it is recognised in equity. Current tax is the expected tax payable on the taxable income for the year, using tax rates enacted or substantially enacted at the Statement of Financial Position date, and any adjustment to tax payable in respect of previous years.

Deferred tax is provided using the Statement of Financial Position liability method, providing for temporary differences between the carrying amounts of assets and liabilities for financial reporting purposes and the amounts used for taxation purposes. The amount of deferred tax provided is based on the expected manner of realisation or settlement of the carrying amount of assets and liabilities, using tax rates enacted or substantially enacted at the Statement of Financial Position date and are expected to apply when the deferred tax asset is realised or the deferred tax liability is settled. A deferred tax asset is recognised only to the extent that it is probable that future taxable profits will be available against which the asset can be utilised. Deferred tax assets are reduced to the extent that it is no longer probable that the related tax benefit will be realised.

Tax consolidation

The Company is the head entity of the tax consolidated group comprising all the Australian wholly owned subsidiaries. The implementation date for the tax consolidated group was 31 May 2006, Current tax expense/income, deferred tax liabilities and deferred tax assets arising from temporary differences of the members of the tax consolidated group are recognised in the separate financial statements of the members of the tax consolidated group using a "stand alone basis without adjusting for intercompany transactions" approach by reference to the carrying amounts of assets and liabilities in the separate financial statements of each entity and the tax values applying under consolidation.

Any current tax Australian liabilities (or assets) and deferred tax assets arising from unused tax losses of the subsidiaries is assumed by the head entity in the tax consolidated group and are recognised as amounts payable (receivable) to (from) other entities in the tax consolidated group. Any difference between these amounts is recognised by the head entity as an equity contribution or distribution.

The Company recognises deferred tax assets arising from unused tax losses of the tax consolidated group to the extent that it is probable that future taxable profits of the tax consolidated group will be available against which the asset can be utilised.

Any subsequent period adjustments to deferred tax assets arising from unused tax losses as a result of revised assessments of the probability of recoverability is recognised by the head entity only.

f) Property, plant and equipment

Plant and equipment is measured at cost less accumulated depreciation and impairment losses.

The cost of fixed assets constructed within the economic entity includes the cost of materials, direct labour, borrowing costs and an appropriate proportion of fixed and variable overheads. Subsequent costs are included in the asset's carrying amount or recognised as a separate asset, as appropriate, only when it is probable that future economic benefits associated with the item will flow to the group and the cost of the item can be reliably. All other repairs and maintenance are charged to the Statement of Comprehensive Income during the financial period in which they are incurred.

Certain expenditure in establishing and commissioning Cyclopharm's PET central Pharmacies has been capitalised. No amortisation has been applied as the asset is not yet deemed held for use. No impairment provision has been deemed appropriate. The Directors are satisfied that the future economic benefits will eventuate to justify the capitalisation of the expenditure incurred.

Impairment

The carrying amount of plant and equipment is reviewed annually by Directors to consider impairment. The recoverable amount is assessed on the basis of the expected net cash flows that will be received from the assets employment and subsequent disposal. The expected net cash flows have been discounted to their present values in determining recoverable amounts.

Depreciation

The depreciable amount of all fixed assets including capitalised lease assets are depreciated on a straight-line basis over their useful lives commencing from the time the asset is held ready for use. Leasehold improvements are depreciated over the shorter of either the unexpired period of the lease or the estimated useful lives of the improvements.

Depreciation is calculated on a straight-line basis over the estimated useful life of the asset as follows:

	Basis	Method
Plant and equipment	10-33%	Straight-line method
Leasehold Improvements	20-50%	Straight-line method
Motor vehicles	20-25%	Straight-line method
	New Patents and licences	Technegas Development costs
Useful lives	Patents - Finite Licenses - Infinite	Finite
Method used	8 - 10 years - Straight line	8 - 10 years - Straight line
Internally generated / Acquired	Acquired	Internally generated
Impairment test / Recoverable Amount testing	Annually and where an indicator of impairment exists	Amortisation method reviewed at each financial year-end; Reviewed annually for indicator of impairment

An item of property, plant and equipment is derecognised upon disposal or when no future economic benefits are expected to arise from the continued use of the asset. Any gain or loss arising on derecognition of the asset (calculated as the difference between the net disposal proceeds and the carrying amount of the item) is included in the Statement of Comprehensive Income in the year the item is derecognised.

g) Borrowing costs

Borrowing costs that are directly attributable to the acquisition, construction or production of assets that necessarily take a substantial period of time to prepare for their intended use or sale, are added to the cost of those assets, until such time as the assets are substantially ready for their intended use or sale.

All other borrowing costs are recognised in the Statement of Comprehensive Income in the year in which they are incurred.

Borrowing costs include interest, amortisation of discounts or premiums relating to borrowings, amortisation of ancillary costs incurred in connection with arrangement of borrowings, foreign exchange losses net of hedged amounts on borrowings, including trade creditors and lease finance charges.

h) Intangibles

Intangible assets

Intangible assets acquired separately are capitalised at cost and from a business combination are capitalised at fair value as at the date of acquisition. Following initial recognition, the cost model is applied to the class of intangible assets.

The useful lives of these intangible assets are assessed to be either finite or indefinite. Where amortisation is charged on assets with finite lives, this expense is taken to the Statement of Comprehensive Income through the 'depreciation and amortisation' line item.

Intangible assets, excluding development costs, created within the business are not capitalised and expenditure is charged against profits in the year in which the expenditure is incurred.

Intangible assets are tested for impairment where an indicator of impairment exists, and in the case of indefinite life intangibles, annually, either individually or at the cash generating unit level. Useful lives are also examined on an annual basis and adjustments, where applicable, are made on a prospective basis.

Research and development costs

Expenditure on research activities is recognised as an expense when incurred.

Expenditure on development activities is capitalised only when it is probable that future benefits will exceed deferred costs and these benefits can be reliably measured. Capitalised development expenditure is stated at cost less accumulated amortisation. Amortisation is calculated using a straight-line method to allocate the costs over a period during which the related benefits are expected to be realised.

Expenditure on the development of the TechnegasPlus generator has been capitalised. A useful life of 9 years has been applied and amortisation for the year included in the Statement of Comprehensive Income. No impairment provision has been deemed appropriate. The Directors are satisfied that the future economic benefits will eventuate to justify the capitalisation of the expenditure incurred.

Expenditure on costs incurred in the application to the Food & Drug Administration authority have been capitalised. A useful life has not been determined as Cyclopharm have not yet received approval from the Food & Drug Administration authority. During the year, \$691,705 was written off the capitalised costs following the settlement of a claim against Clinquest Inc. No impairment provision has been deemed necessary at balance date. The Directors are satisfied that the future economic benefits will eventuate to justify the carrying value of the capitalised expenditure.

Development expenditure is tested annually for impairment or more frequently if events or changes in circumstances indicate that it might be impaired. Capitalised development expenditure is measured at cost less any accumulated amortisation and impairment losses.

i) Inventories

Inventories are valued at the lower of cost and net realisable value where net realisable value is the estimated selling price in the ordinary course of business, less estimated costs of completion and the estimated costs necessary to make the sale.

Costs incurred in bringing each product to its present location and conditions are accounted for as follows:

- Raw materials: purchase cost on a first-in, first-out basis;
- Finished goods and work-in-progress: cost of direct materials and labour and an appropriate portion of manufacturing overheads based on normal operating capacity but excluding borrowing costs.

j) Trade and other receivables

Trade receivables, which generally have 30-90 day terms, are recognised and carried at original invoice amount less an allowance for any uncollectible amounts. A specific estimate for doubtful debts is made when collection of the full amount is no longer probable. Bad debts are written off when identified.

k) Cash and cash equivalents

Cash and cash equivalents comprise cash on hand, deposits held at call with banks, short-term deposits with an original maturity of three months or less and bank overdrafts. Bank overdrafts are

shown within short-term borrowings in current liabilities on the Statement of Financial Position. For the purposes of the Statement of Cash Flows, cash and cash equivalents consist of cash and cash equivalents as defined above, net of outstanding bank overdrafts.

l) Trade and other payables

Trade payables and other payables are carried at amortised costs and represent liabilities for goods and services provided to the Group prior to the end of the financial year that are unpaid and arise when the Group becomes obliged to make future payments in respect of the purchase of these goods and services. Trade payables are normally settled within 30 to 60 days.

m) Interest-bearing loans and borrowings

All loans and borrowings are initially recognised at cost, being the fair value of the consideration received net of issue costs associated with the borrowing. After initial recognition, interest-bearing loans and borrowings are subsequently measured at amortised cost using the effective interest method. Amortised cost is calculated by taking into account any issue costs and any discount or premium on settlement. Gains and losses are recognised in the Statement of Comprehensive Income when the liabilities are derecognised and as well as through the amortisation process.

n) Provisions

Provisions are recognised when the Group has a present obligation (legal or constructive) as a result of past events, for which it is probable that an outflow of economic benefits will result and that an outflow can be reliably measured. Where the Group expects some or all of a provision to be reimbursed, for example under an insurance contract, the reimbursement is recognised as a separate asset but only when the reimbursement is virtually certain. The expense relating to any provision is presented in the Statement of Comprehensive Income net of any reimbursement.

o) Employee entitlements

Provision is made for employee benefits accumulated as a result of employees rendering services up to the reporting date. These benefits include wages and salaries, annual leave and long service leave.

Employee benefits expected to be settled within twelve months of the reporting date are measured at their nominal amounts based on remuneration rates which are expected to be paid when the liability is settled plus related on-costs. All other employee benefit liabilities are measured at the present value of the estimated future cash outflow to be made in respect of services provided by employees up to the reporting date. In determining the present value of future cash outflows, the market yield as at the reporting date on national government bonds, which have terms to maturity approximating the terms of the related liability, are used.

Employee benefit expenses and revenues arising in respect of wages and salaries, non-monetary benefits, annual leave, long service leave and other leave benefits; and other types of employee benefits are recognised against profits on a net basis in their respective categories.

p) Employee share and performance share schemes

The fair value of performance rights issued under the Cyclopharm Long Term Incentive Plan are recognised as a personnel expense over the vesting period with a corresponding increase in Employee Equity Benefits Reserve.

The fair value of performance the implied option attached to shares granted is determined using a pricing model that takes into account factors that include exercise price, the term of the performance option, the vesting and performance criteria, the share price at grant date and the expected price volatility of the underlying share. The fair value calculation excludes the impact of any non market vesting conditions. Non market vesting conditions are included in assumptions about the number of performance options that are expected to become exercisable. At each balance date, the entity revises its estimate of the number of performance rights that are expected to become exercisable. The personnel expense recognised each period takes into account the most recent estimate.

Shares issued under employee and executive share plans are held in trust until vesting date. Unvested shares held by the trust are consolidated into the group financial statements.

q) Leases

Operating Leases

Leases where the lessor retains substantially all the risks and benefits of ownership of the asset are classified as operating leases. Operating lease payments are recognised as an expense in the Statement of Comprehensive Income on a straight-line basis over the lease term. Lease incentives under operating leases are recognised as a liability and amortised on a straight-line basis over the life of the lease.

r) Revenue recognition

Revenue is recognised to the extent that it is probable that the economic benefits will flow to the Group and the revenue can be reliably measured. The following specific recognition criteria must also be met before revenue is recognised:

Sale of goods

Revenue is recognised (net of returns, discounts and allowances) when the significant risks and rewards of ownership and therefore control of the goods have passed to the buyer and can be measured reliably. Control is considered to have passed to the buyer at the time of delivery of the goods to the customer.

Interest

Revenue is recognised as the interest accrues using the effective interest rate method, which is the rate that exactly discounts estimated future cash receipts through the expected life of the financial instrument to the net carrying amount of the financial asset.

Dividends

Revenue from quoted investments is recognised in the Statement of Comprehensive Income on the day which the relevant investment is first quoted on an "ex-basis". Dividend revenue is recognised net of any franking credits.

Revenue from distributions from controlled entities is recognised by the Company when they are declared by the controlled entities. Revenue from dividends from associates and other investments is recognised when dividends are received. Dividends received out of pre-acquisition reserves are eliminated against the carrying amount of the investment and not recognised in revenue.

Research and development grants

Where a grant is received relating to research and development costs that have been expensed, the grant is recognised as revenue.

All revenue is stated net of the amount of goods and services tax ("GST").

s) Other taxes

Revenues, expenses and assets are recognised net of the amount of GST except where the GST incurred is not recoverable from the Australian Taxation Office ("ATO"), and is therefore recognised as part of the asset's cost or as part of the expense item. Receivables and payables are stated inclusive of GST. The net amount of GST recoverable from, or payable to, the ATO is included as part of receivables or payables in the Statement of Financial Position. Cash flows are presented in the Statement of Cash Flows on a gross basis and the GST component of cash flows arising from investing and financing activities, which is recoverable from, or payable to the taxation authority are classified as operating cash flows.

t) Financial instruments

Financial instruments are initially measured at cost on trade date, which includes transaction costs, when the related contractual rights or obligations exist. Subsequent to initial recognition these instruments are measured as set out below.

Financial assets at fair value through profit and loss

A financial asset is classified in this category if acquired principally for the purpose of selling in the short term, or if so designated by management and within the requirement of AASB139: Recognition and Measurement of Financial Instruments.

Loans and receivables

Loans and receivables are non-derivative financial assets with fixed or determinable payments that are not quoted in an active market and are stated at amortised cost using the effective interest rate method.

De-recognition of financial instruments

Financial assets

A financial asset (or, where applicable, a part of a financial asset or part of a group of similar financial assets) is derecognised when:

- the rights to receive cash flows from the asset have expired;
- the Group retains the right to receive cash flows from the asset, but has assumed an obligation to pay them in full without material delay to a third party under a 'pass-through' arrangement; or
- the Group has transferred its rights to receive cash flows from the asset and either (a) has transferred substantially all the risks and rewards of the asset, or (b) has neither transferred nor retained substantially all the risks and rewards of the asset, but has transferred control of the asset.

When the Group has transferred its rights to receive cash flows from an asset and has neither transferred nor retained substantially all the risks and rewards of the asset nor transferred control of the asset, the asset is recognised to the extent of the Group's continuing involvement in the asset. Continuing involvement that takes the form of a guarantee over the transferred asset is measured at the lower of the original carrying amount of the asset and the maximum amount of consideration received that the Group could be required to repay.

Financial liabilities

A financial liability is derecognised when the obligation under the liability is discharged or cancelled or expires. When an existing financial liability is replaced by another from the same lender on substantially different terms, or the terms of an existing liability are substantially modified, such an exchange or modification is treated as a de-recognition of the original liability and the recognition of a new liability, and the difference in the respective carrying amounts is recognised in profit or loss.

Impairment of financial assets

The Group assesses at each Statement of Financial Position date whether a financial asset or group of financial assets is impaired.

Financial assets carried at amortised cost

If there is objective evidence that an impairment loss on loans and receivables carried at amortised cost has been incurred, the amount of the loss is measured as the difference between the asset's carrying amount and the present value of estimated future cash flows (excluding future credit losses that have not been incurred) discounted at the financial asset's original effective interest rate (i.e. the effective interest rate computed at initial recognition). The carrying amount of the asset is reduced either directly or through use of an allowance account. The amount of the loss is recognised in profit or loss.

The Group first assesses whether objective evidence of impairment exists individually for financial assets that are individually significant, and individually or collectively for financial assets that are not individually significant. If it is determined that no objective evidence of impairment exists for an individually assessed financial asset, whether significant or not, the asset is included in a group of financial assets with similar credit risk characteristics and that group of financial assets is collectively assessed for impairment. Assets that are individually assessed for impairment and for which an

impairment loss is or continues to be recognised are not included in a collective assessment of impairment.

If, in a subsequent period, the amount of the impairment loss decreases and the decrease can be related objectively to an event occurring after the impairment was recognised, the previously recognised impairment loss is reversed. Any subsequent reversal of an impairment loss is recognised in profit or loss, to the extent that the carrying value of the asset does not exceed its amortised cost at the reversal date.

Financial assets carried at cost

If there is objective evidence that an impairment loss has been incurred on an unquoted equity instrument that is not carried at fair value (because its fair value cannot be reliably measured), or on a derivative asset that is linked to and must be settled by delivery of such an unquoted equity instrument, the amount of the loss is measured as the difference between the asset's carrying amount and the present value of estimated future cash flows, discounted at the current market rate of return for a similar financial asset.

u) Contributed equity

Share capital

Ordinary shares are classified as equity. Incremental costs directly attributable to the issue of new shares or options are shown in equity as a deduction, net of tax, from the proceeds.

Other contributed equity

In accordance with *UIG 1052 Tax Consolidation Accounting*, additional contributed equity was recorded to recognise the transfer of tax liabilities from Vita Medical Limited to Vita Life Sciences Limited, being the parent of the Australian tax consolidated group at the relevant time. This event occurred prior to Cyclopharm acquiring its interests in the net assets of Vita Medical Limited.

As part of the restructure a subsidiary of Cyclopharm, Vita Medical Australia Pty Ltd acquired all the assets, liabilities and business from Vita Medical Limited, the former group parent.

With effect from 31 May 2006, Cyclopharm also acquired 100% of the other group operating subsidiaries from the ultimate holding company, Vita Life Sciences Limited. Accordingly, the group comprises Cyclopharm and the following wholly owned subsidiaries:

- Cyclomedica Australia Pty Ltd (formerly Vita Medical Australia Pty Ltd)
- Cyclomedica Ireland Ltd (formerly Vitamedica Europe Ltd)
- Cyclomedica Europe Ltd
- Cyclomedica Canada Limited (formerly Vita Medical Canada Ltd)
- Cyclomedica Germany GmbH
- Allrad 28 Pty Ltd
- Allrad 29 Pty Ltd

These entities collectively comprise the medical diagnostic equipment and associated consumables business formerly operated as the Vita Medical Group – now known as the Cyclopharm Group. The transaction has been accounted for as a 'reverse acquisition' as defined in *AASB 3 Business Combinations* whereby Cyclopharm is the legal parent and Cyclomedica Australia Pty Limited is the financial parent, which for accounting purposes is deemed to be the acquirer.

The consideration for the minority interests of the controlled entities and costs of acquisition have been charged to other contributed equity in accordance with *AASB 127 Consolidated and Separate Financial Statements*.

v) Earnings per share

Basic earnings per share

Basic earnings per share is determined by dividing the net profit/(loss) after income tax attributable to members of the Company by the weighted average number of ordinary shares outstanding during the financial year. Where a change in the number of ordinary shares on issue without a corresponding change in recognised resources during the year the number of ordinary shares for all periods presented are correspondingly adjusted as if the event had occurred at the beginning of the earliest period presented.

Diluted earnings per share

Diluted earnings per share adjusts the figures used in the determination of basic earnings per share to take into account the after income tax effect of interest and other financing costs associated with dilutive potential ordinary shares and the weighted average number of shares assumed to have been issued for no consideration in relation to dilutive potential ordinary shares. Where a change in the number of ordinary shares on issue without a corresponding change in recognised resources during the year the number of ordinary shares for all periods presented are correspondingly adjusted as if the event had occurred at the beginning of the earliest period presented.

3. SEGMENT REPORTING

The Group's primary segment reporting format is business segments as the Group's risks and returns are affected predominantly by differences in the products and services produced. The Group's secondary segment is geographical.

The operating businesses are organised and managed separately according to the nature of the products and services provided, with each segment representing a strategic business unit that offers different products and serves different markets.

The Technegas segment is a supplier of diagnostic equipment and consumables used by physicians in the detection of pulmonary embolism.

The Molecular Imaging segment will produce radiopharmaceuticals to be used by physicians in the detection of cancer, neurological disorders and cardiac disease.

Transfer prices between business segments are set on an arm's length basis in a manner similar to transactions with third parties. Segment revenue, segment expense and segment result include transfers between business segments. Those transfers are eliminated on consolidation.

Business segments

The tables under the heading business segments present revenue and profit information and certain asset and liability information regarding business segments for the years ended 31 December 2010 and 31 December 2009.

Geographical segments

The tables under the heading geographical segment present revenue and profit information and certain asset and liability information regarding geographical segments for the years ended 31 December 2010 and 31 December 2009.

Notes

Continued

3. SEGMENT REPORTING (continued)

Business Segments

For the period ended	Consolidated		
	Technegas	Molecular Imaging	Total
31 December 2010	\$	\$	\$
Revenue			
Sales to external customers	9,442,763	21,842	9,464,605
Finance revenue	63,802	5,170	68,972
Total revenue	9,506,565	27,012	9,533,577
Result			
Profit / (loss) before tax and finance costs	1,845,650	(838,943)	1,006,707
Finance costs	(172,857)	(25,442)	(198,299)
Profit / (Loss) before income tax	1,672,793	(864,385)	808,408
Income tax expense	(358,302)	-	(358,302)
Profit / (Loss) after income tax	1,314,491	(864,385)	450,106
Assets and liabilities			
Segment assets	10,603,331	11,538,568	22,141,899
Segment liabilities	(3,152,832)	(5,222,913)	(8,375,745)
Other segment information			
Depreciation and amortisation	(317,072)	(33,474)	(350,546)

Notes

Continued

3. SEGMENT REPORTING (continued)

Business Segments

For the period ended	Consolidated		
	Technegas	Molecular Imaging	Total
31 December 2009	\$	\$	\$
Revenue			
Sales to external customers	11,098,579	-	11,098,579
Finance revenue	22,285	70,901	93,186
Other revenue	2,403,090	-	2,403,090
Total revenue	13,523,954	70,901	13,594,855
Result			
Profit / (loss) before tax and finance costs	2,955,015	(250,011)	2,705,004
Finance costs	(136,759)	(744)	(137,503)
Profit / (Loss) before income tax	2,818,256	(250,755)	2,567,501
Income tax expense	(523,011)	-	(523,011)
Profit / (Loss) after income tax	2,295,245	(250,755)	2,044,490
Assets and liabilities			
Segment assets	13,452,842	6,191,569	19,644,411
Segment liabilities	(3,583,244)	(2,132,526)	(5,715,770)
Other segment information			
Impairment write downs	(691,705)	-	(691,705)
Depreciation and amortisation	(450,229)	-	(450,229)

Notes

Continued

Geographical Segments

For the year ended	Asia Pacific	Europe	Canada	Other	Total
31 December 2010	\$	\$	\$	\$	\$
Revenue					
Sales to external customers	2,280,115	5,487,463	1,598,679	98,348	9,464,605
Finance revenue	59,832	9,140	-	-	68,972
Total segment revenue	2,339,947	5,496,603	1,598,679	98,348	9,533,577
Assets and liabilities					
Segment assets	17,282,538	4,075,722	783,639	-	22,141,899
Segment liabilities	(7,985,879)	(255,157)	(134,709)	-	(8,375,745)

For the year ended	Consolidated				Total
	Asia Pacific	Europe	Canada	Other	
31 December 2009	\$	\$	\$	\$	\$
Revenue					
Sales to external customers	1,886,030	7,309,497	1,734,492	168,560	11,098,579
Finance revenue	89,994	3,192	-	-	93,186
Other revenue	2,403,090	-	-	-	2,403,090
Total segment revenue	4,379,114	7,312,689	1,734,492	168,560	13,594,855
Assets and liabilities					
Segment assets	13,858,482	5,210,112	575,817	-	19,644,411
Segment liabilities	(5,082,328)	(531,659)	(101,783)	-	(5,715,770)

4. REVENUES AND EXPENSES

		Consolidated	
		2010	2009
Notes		\$	\$
Revenue			
	Sales revenue	9,464,605	11,098,579
	Finance revenue	68,972	93,186
Other Revenue			
	Realised foreign exchange gains	-	389,249
	Clinquest settlement proceeds	-	2,013,841
	Total other revenue	-	2,403,090
Expenses			
a) Cost of materials and manufacturing			
	Cost of materials and manufacturing	2,308,992	2,546,484
b) Finance costs			
	Interest paid on loans from external parties	198,299	137,503
c) Depreciation and amortisation			
	Depreciation of plant and equipment	291,220	398,671
	Depreciation of leasehold improvements	10,914	6,241
	Amortisation of intangibles	48,412	45,317
		350,546	450,229
d) Research & development			
	Research and development expense	16,847	39,728
		16,847	39,728
e) Employee benefits expense			
	Salaries and wages	2,698,471	3,307,153
	Non-Executive Director fees and consultant costs	162,583	156,548
	Share-based payments expense	96,177	74,283
23a		2,957,231	3,537,984
f) Administration expense			
	Legal and professional costs	543,391	1,581,287
	Office and facility costs	662,997	736,873
	Travel and motor vehicle costs	457,115	557,085
		1,663,503	2,875,245
g) Other expenses			
	Writedown of capitalised FDA development cost	-	691,705
	Other	337,067	92,629
		337,067	784,334

5. INCOME TAX

	Consolidated	
	2010	2009
	\$	\$
Current income tax (expense) / benefit	(242,522)	(563,249)
Deferred tax (expense) / benefit	(115,780)	40,238
Income tax reported in income statement	(358,302)	(523,011)

A reconciliation of income tax benefit / (expense) applicable to accounting profit before income tax at the statutory income tax rate to income tax expense at the Group's effective income tax rate is as follows:

Accounting profit / (loss) before income tax	808,408	2,567,501
Statutory income tax rate of 30%	(242,522)	(770,250)
Expenditure not allowable for income tax purposes	(1,208)	(1,218)
Share based payments for which no deduction is obtained	(28,853)	(22,285)
Deferred expenditure	(146,089)	-
Effects of lower rates on overseas income	39,254	207,001
Tax losses not recognised in foreign subsidiaries	21,116	63,741
Total income tax (expense) / benefit	(358,302)	(523,011)
Effective income tax rate	(44.3%)	(20.4%)
Current tax liabilities		
Current income tax liability	318,850	174,039
Deferred tax assets/liabilities		
Deferred tax assets and liabilities relate to the following:		
Deferred tax assets from temporary differences on:		
Provisions	223,418	154,629
Tax losses applied in current year	(475,955)	(447,450)
Tax losses of parent entity brought to account	413,273	384,768
Tax losses / (payable) transferred from Australian subsidiaries	62,682	(117,850)
Adjustment to recognise deferred tax asset arising from share issue costs	48,389	25,901
Transfer to deferred tax liability	(365,219)	-
Investment accounted for using equity method	93,412	-
Other	-	2
Total deferred tax assets	-	-
Deferred tax liabilities from temporary differences on:		
Capitalised expenditure	709,876	367,304
Transfer to deferred tax assets	(365,219)	-
Total deferred tax liabilities	344,657	367,304

Notes

Continued

6. NET TANGIBLE ASSETS AND EARNINGS PER SHARE

Net Tangible Assets per share

	Consolidated	
	2010	2009
	\$	\$
Net assets per share	0.08	0.08
Net tangible assets per share	0.07	0.07
	Number	Number
Weighted average number of ordinary shares for net assets per share	171,012,616	171,012,616
	2010	2010
	\$	\$
Net assets	13,766,154	13,928,641
Net tangible assets	11,145,002	11,506,974

Earnings per share

	Consolidated	
	2010	2009
	cents	cents
Basic earnings per share for continuing operations	0.26	1.20
Basic earnings per share	0.26	1.20
Diluted earnings per share	0.26	1.20
	Number	Number
Weighted average number of ordinary shares for basic and diluted earnings per share	171,012,616	171,012,616
	2010	2010
	\$	\$
Earnings used to calculate basic earnings per share	450,106	2,044,490
Earnings used to calculate diluted earnings per share	450,106	2,044,490

7. CASH AND CASH EQUIVALENTS

	Consolidated	
	2010	2009
	\$	\$
Cash at bank and in hand	1,541,644	4,612,205
Total cash and cash equivalents	1,541,644	4,612,205

Cash at bank and in hand earns interest at floating rates based on daily bank deposit rates.

The fair value of cash equivalents is \$1,541,644 (2009: \$4,612,205).

Reconciliation of Cash Flow Statement

For the purpose of the Cash Flow Statement, cash and cash equivalents comprise the following:

	2010	2009
Cash at bank and in hand	1,541,644	4,612,205
	1,541,644	4,612,205

(a) Reconciliation of net profit / (loss) after tax to net cash flows from operations

	2010	2009
Net profit / (loss) after tax	450,106	2,044,490
Adjustments for non-cash income and expense items:		
Depreciation	302,134	404,912
Amortisation	48,412	45,317
Impairment written down	-	(691,705)
Movement in equity	-	325,957
Movement in investment in associate	311,769	-
Movement provision for employee benefits	64,078	113,522
Movement in foreign exchange	(577,774)	-
Movement in employee benefits reserve	96,177	74,283
Movement in other provisions	(78,109)	24,609
	616,793	2,341,385
Increase/decrease in assets and liabilities:		
(Increase) / decrease in receivables	71,626	812,099
(Increase) / decrease in inventories	(5,875)	(387,626)
(Increase) / decrease in other receivables	(78,007)	255,250
(Increase) / decrease in deferred tax assets	-	616,379
Increase / (decrease) in creditors	(817,661)	339,130
Increase / (decrease) in current tax liabilities	144,811	168,968
Increase / (decrease) in deferred tax liabilities	(22,647)	(454,552)
Net cash from operating activities	(90,960)	3,691,033

8. TRADE AND OTHER RECEIVABLES

		Consolidated	
		2010	2009
Notes		\$	\$
Current			
	Trade receivables, third parties	3,680,579	3,590,705
	Other receivables	522,595	690,664
5	Current income tax receivable	92,313	-
Total Current trade and other receivables		4,295,487	4,281,369
Non-current			
	Other receivables	161,500	-
Total Non-current other receivables		161,500	-
Total Current trade and other receivables		4,456,987	4,281,369

Terms and conditions

Terms and conditions relating to the above financial instruments

- a. Trade receivables are non-interest bearing and generally on 30 and 60 day terms.
- b. Other debtors are non-interest bearing and have repayment terms between 30 and 90 days.
- c. Related party details are set out in the Note 19 Related party disclosures, controlled entities.

9. INVENTORIES

		Consolidated	
		2010	2009
Notes		\$	\$
Current			
	Raw materials at cost	625,955	853,415
	Finished goods at low er of cost or net realisable value	1,920,912	2,389,577
Total current inventory		2,546,867	3,242,992
Non-current			
	Finished goods at low er of cost or net realisable value	702,000	-
Total non-current inventory		702,000	-
Total inventory		3,248,867	3,242,992

10. PROPERTY, PLANT AND EQUIPMENT

Year ended						
31 December 2010	Leasehold Land and buildings	Leasehold improvements	Plant and equipment	Leased Plant and Equipment	Capital Work in Progress	Total
Consolidated	\$	\$	\$	\$	\$	\$
1 January 2010						
at written down value	161,500	19,631	726,433	-	4,145,387	5,052,951
Additions / Transfers	1,983,729	2,825,047	5,091,490	26,962	-	9,927,228
Disposals / Transfers	(161,500)	-	(324,667)	-	(4,132,350)	(4,618,517)
Depreciation for the year	(3,261)	(10,914)	(287,960)	-	-	(302,134)
31 December 2010						
at written down value	1,980,468	2,833,764	5,205,297	26,962	13,037	10,059,528
1 January 2010						
Cost value	161,500	208,116	2,527,570	114,049	4,145,387	7,156,622
Accumulated depreciation	-	(188,485)	(1,801,137)	(114,049)	-	(2,103,671)
Net carrying amount	161,500	19,631	726,433	-	4,145,387	5,052,951
31 December 2010						
Cost value	1,983,729	3,033,163	7,053,894	141,011	13,037	12,224,834
Accumulated depreciation	(3,261)	(199,400)	(1,848,598)	(114,049)	-	(2,165,307)
Net carrying amount	1,980,468	2,833,764	5,205,297	26,962	13,037	10,059,528

During the year, the PET Facility at Macquarie University Hospital, New South Wales was commissioned and \$4,132,350 of Capital Work in Progress was transferred to the Leasehold Land and Buildings, Leasehold improvements and the Plant and Equipment asset classes. The remaining Capital Work in Progress balance of \$13,037 relates solely to the development of the PET Facility at Kensington, Victoria.

Year ended						
31 December 2009	Land and buildings	Leasehold improvements	Plant and equipment	Leased Plant and Equipment	Capital Work in Progress	Total
Consolidated		\$	\$	\$	\$	\$
1 January 2009						
at written down value	161,500	23,945	1,165,043	-	1,375,346	2,725,834
Additions	-	1,927	134,135	-	2,770,041	2,906,103
Disposals / Transfers	-	-	(174,074)	-	-	(174,074)
Depreciation for the year	-	(6,241)	(398,671)	-	-	(404,912)
31 December 2009						
at written down value	161,500	19,631	726,433	-	4,145,387	5,052,951
1 January 2009						
Cost value	161,500	206,189	2,796,499	114,049	1,375,346	4,653,583
Accumulated depreciation	-	(182,244)	(1,631,456)	(114,049)	-	(1,927,749)
Net carrying amount	161,500	23,945	1,165,043	-	1,375,346	2,725,834
31 December 2009						
Cost value	161,500	208,116	2,527,570	114,049	4,145,387	7,156,622
Accumulated depreciation	-	(188,485)	(1,801,137)	(114,049)	-	(2,103,671)
Net carrying amount	161,500	19,631	726,433	-	4,145,387	5,052,951

11. INVESTMENTS IN ASSOCIATE

				Consolidated	
				2010	2009
				\$	\$
Associated companies				188,231	-
Name	Principal Activities	Country of Incorporation	Shares	Ownership Interest	
				2010	2009
Macquarie Medical Imaging Pty Ltd	Imaging centre	Australia	Preference	20%	0%
				Consolidated	
				2010	2009
				\$	\$
Macquarie Medical Imaging Pty Ltd				-	-
At 1 January				-	-
Investment in associate				500,000	-
Sales to associate				(397)	-
Share of losses after income tax				(311,372)	-
At 31 December				188,231	-
				Consolidated	
				2010	2009
				\$	\$
Extract from the associates statement of financial position:					
Current Assets				2,100,522	
Non-current Assets				4,108,207	-
Current Liabilities				1,413,137	-
Non-current Liabilities				3,652,567	-
Net assets				1,143,025	-
Share of associates' net assets				228,605	
				Consolidated	
				2010	2009
				\$	\$
Extract from the associates statement of comprehensive income:					
Revenue				1,313,215	-
Net Loss				(1,556,857)	-

Cyclopharm's wholly owned subsidiary Cyclopet Pty Ltd has invested \$500,000 in Macquarie Medical Imaging Pty Ltd, an imaging joint venture at Macquarie University Hospital. Cyclopet Pty Ltd has a 20% interest in Macquarie Medical Imaging Pty Ltd. The share of the associate's loss for the period was \$311,372. Commercial operations commenced in July 2010.

The fair value of the Group's investment in Macquarie Medical Imaging Pty Ltd is \$188,231.

11. INVESTMENTS IN ASSOCIATE (continued)

	Consolidated	
	2010	2009
	\$	\$
Contingent liabilities relating to the associates:		
Guarantee of associates financing facilities for which the Group is severally liable	16,000,000	-

The guarantee by Cyclopharm Limited and Cyclopet Pty Limited is security for the whole Macquarie Medical Imaging Pty Ltd financing facility provided by the Commonwealth Bank of Australia. Cyclopharm Group's liability is limited to the amount that Cyclopharm Limited and Cyclopet Pty Limited are obliged to fund under a Subscription Agreement with Alfred Health Solutions Pty Ltd which at the date of this report is 20% of the liability.

12. INTANGIBLE ASSETS

	Intellectual property	Technegas Development	FDA Development	Total
	\$	\$	\$	\$
Consolidated				
Balance at				
1 January 2009	110,083	161,916	2,149,668	2,421,667
Arising during the year	37,311	-	210,586	247,897
Amortisation	(17,692)	(30,720)	-	(48,412)
Balance at				
31 December 2010	129,702	131,196	2,360,254	2,621,152
31 December 2010				
Non-Current	129,702	131,196	2,360,254	2,621,152
Total	129,702	131,196	2,360,254	2,621,152
31 December 2009				
Non-Current	110,083	161,916	2,149,668	2,421,667
Total	110,083	161,916	2,149,668	2,421,667

The recoverable amount of FDA and Technegas development costs have been assessed using a discounted cash flow methodology forecasting six years of pre-tax cash flows.

The following describes each key assumption on which management has based its value in use calculations:

- Six year pre- tax cash flow projections, based upon management approved budgets and growth rates covering a one year period, with the subsequent periods based upon management expectations of growth excluding the impact of possible future acquisitions, business improvement capital expenditure and restructuring.
- The discount factor used was 15% in 2009 (2009: 15%).
- The Directors have concluded that the recoverable amount of the FDA development costs and other intangibles exceed their carrying value.
- In February 2011, the FDA approved the Company's Special Protocol Assessment. The SPA ensures that the agreed clinical trial design meets the FDA's expectations for a pivotal study. Cyclopharm's positive SPA response leaves no clinical issues outstanding, and provides Cyclopharm with confidence that the design of the Phase III development program and clinical trial for Technegas is suitable to support regulatory approval for the United States.

Notes

Continued

13. TRADE AND OTHER PAYABLES

	Consolidated	
	2010	2009
	\$	\$
Trade payables, third parties	688,933	1,414,254
Other payables and accruals	393,559	485,899
Total trade and other payables	1,082,492	1,900,153

Terms and conditions

Terms and conditions relating to the above financial instruments:

- (i) Trade payables are non-interest bearing and are normally settled on 30-60 day terms.
- (ii) Other payables and accruals are non-interest bearing and have an average term of 4 months.
- (iii) The non-interest bearing loan, related party loan is payable when called upon. Related party details are set out in the Note 19 Related party disclosures, controlled entities

14. INTEREST BEARING LOANS AND BORROWINGS

	Consolidated	
	2010	2009
	\$	\$
Current		
Lease liability	2,400	-
Bank loan - secured	6,075,900	1,757,350
Interest bearing loans and borrowings (current)	6,078,300	1,757,350
Non-current		
Lease liability	24,453	-
Bank loan - secured	-	975,900
Interest bearing loans and borrowings (non-current)	24,453	975,900
Total interest bearing loans and borrowings	6,102,753	2,733,250

14. INTEREST BEARING LOANS AND BORROWINGS (continued)

(a) Financing facilities available:

At reporting date, the following financing facilities had been negotiated and were available:

		Consolidated	
		2010	2009
Notes		\$	\$
Total facilities available:			
	- secured bank loans, third party	6,075,900	6,450,000
		6,075,900	6,450,000
Facilities used at reporting date:			
	- secured bank loans, third party	14 6,075,900	2,733,250
		6,075,900	2,733,250
Facilities unused at reporting date:			
	- secured bank loans, third party	-	3,716,750
		-	3,716,750
	Total facilities	6,075,900	6,450,000
	Facilities used at reporting date:	(6,075,900)	(2,733,250)
	Facilities unused at reporting date:	-	3,716,750

(b) Secured Bank Loans

- (i) Cyclopharm has an amortising bank bill facility provided by the National Australia Bank of \$1.35 million. The entirety of the facility must be repaid by 31 July 2011. The facility is secured by a first registered mortgage debenture over Cyclopharm Limited and a guarantee and indemnity for \$6,450,000 from Cyclomedica Australia Pty Ltd, CycloPET Pty Ltd, Allrad No. 28 Pty Ltd and Allrad No. 29 Pty Ltd. Supported by Fixed and Floating Charge and First Registered Debenture charges over these companies.
- (ii) Cyclopharm has a 15 month multi-option facility (MOF) provided by the National Australia Bank for \$5.1 million. The facility expires in March 2011. The facility is secured by a first registered mortgage debenture over Cyclopharm Limited and a guarantee and indemnity for \$6,450,000 from Cyclomedica Australia Pty Ltd, CycloPET Pty Ltd, Allrad No. 28 Pty Ltd and Allrad No. 29 Pty Ltd. Supported by Fixed and Floating Charge and First Registered Debenture charges over these companies.

15. PROVISIONS

	Consolidated		
	Employee Entitlements	Other	Total
Consolidated	\$	\$	\$
Balance at			
<i>1 January 2010</i>	462,915	78,109	541,024
Arising during the year	200,545	-	200,545
Utilised	(136,467)	(78,109)	(214,576)
Balance at			
<i>31 December 2010</i>	526,993	-	526,993
31 December 2010			
Current	499,283	-	499,283
Non-Current	27,710	-	27,710
Total	526,993	-	526,993
Number of employees			
Number of employees at year end	38		
31 December 2009			
Current	420,174	78,109	498,283
Non-Current	42,741	-	42,741
Total	462,915	78,109	541,024
Number of employees			
Number of employees at year end	41		

Notes

Continued

16. CONTRIBUTED EQUITY

		Consolidated			
		2010	2009	2010	2009
Notes	Number	Number	\$	\$	
Issued and paid up capital					
	Ordinary shares	(a) 171,012,616	171,012,616	16,422,066	16,422,066
	Other contributed equity	(b) -	-	(5,333,158)	(5,333,158)
Total issued and paid up capital		171,012,616	171,012,616	11,088,908	11,088,908
Ordinary shares					
(a) Issued and paid up capital					
	Balance at the beginning of the period	171,012,616	171,012,616	16,422,066	16,422,066
	Balance at end of period	171,012,616	171,012,616	16,422,066	16,422,066
(b) Other contributed equity					
	Balance at the beginning of the period	-	-	(5,333,158)	(5,295,657)
	Acquisition of minority interests in controlled entities	-	-	-	(37,501)
	Balance at end of period	-	-	(5,333,158)	(5,333,158)

Ordinary shares have the right to receive dividends as declared and, in the event of winding up the Company, to participate in the proceeds from the sale of all surplus assets in proportion to the number of and amounts paid up on shares held. Ordinary shares entitle their holder to one vote, either in person or by proxy, at a meeting of the Company.

16. CONTRIBUTED EQUITY (continued)

When managing capital, management's objective is to ensure the entity continues as a going concern as well as to maintain optimal returns for shareholders and benefits for other stakeholders. Management also aims to maintain a capital structure that ensures the lowest cost of capital available to the entity.

Management constantly assess the capital structure to take advantage of favourable costs of capital and/or high returns on assets. As the market is continually changing, management may issue dividends to shareholders, issue new shares, increase its short or long term borrowings or sell assets to reduce borrowings.

The Directors did not declare a dividend during the financial year ended 31 December 2010.

Management monitor capital through the gearing ratio (net debt/total capital). Management aim to ensure that the Group's gearing ratio does not exceed 45%. The Group has satisfied its year-end externally imposed capital requirements of its banking facilities detailed in Note 14 (b).

	Notes	Consolidated	
		2010 \$	2009 \$
Total interest bearing loans and borrowings	14	6,102,753	2,733,250
Less cash and cash equivalents	7	(1,541,644)	4,612,205
Net (cash) / debt		4,561,109	(1,878,955)
Total equity		13,766,154	13,928,641
Gearing ratio*		33.1%	(13.5%)

*A negative ratio denotes that net cash exceeded net borrowings at the reporting date

17. FINANCIAL RISK MANAGEMENT OBJECTIVES

The Group's principal financial instruments comprise receivables, payables, bank loans and overdrafts and cash and short-term deposits. The Group manages its exposure to key financial risks, including interest rate and currency risk in accordance with the Group's financial risk management policy. The objective of the policy is to support the delivery of the Group's financial targets while protecting future financial security.

The Group uses different methods to measure and manage different types of risks to which it is exposed. These include monitoring levels of exposure to interest rate, foreign exchange risk and assessments of market forecasts for interest rate, foreign exchange and commodity prices. Ageing analysis and monitoring of specified credit allowances are undertaken to manage credit risk, liquidity risk is monitored through the development of future rolling cash flow forecasts.

The Board review and agrees policies for managing each of these risks as summarised below.

Primary responsibility for identification and control of financial risks rests with the Audit and Risk Management Committee under the authority from the Board. The Board reviews and agrees policies for managing each of the risks identified below, including for interest rate risk, credit allowances and cash flow forecast projections. It is, and has been throughout the period under review, the Group's policy that no trading in financial instruments shall be undertaken.

Details of the significant accounting policies and methods adopted, including the criteria for recognition, the basis of measurement and the basis on which income and expenses are recognised, in respect of each class of financial asset, financial liability and equity instrument are disclosed in Note 2.

Notes

Continued

(a) Interest rate risk

The Group's exposure to the risk of changes in market interest rates relates primarily to the Group's long term debt obligations with a floating interest rate. The Group's policy is to manage its interest cost using a mix of fixed and variable rate debt. The Group constantly analyses its interest rate exposure. Within this analysis consideration is given to potential renewals of existing positions, alternative financing, positions and the mix of fixed and variable interest rates.

The following sensitivity analysis is based on the interest rate risk exposures in existence at the Statement of Financial Position date.

At 31 December 2010, if interest rates had moved, as illustrated in the table below, with all other variables held constant, pre tax profit would have been affected as follows:

Notes	Consolidated	
	2010 \$	2009 \$
Judgements of reasonably possible movements:		
Profit / (loss) before income tax		
+1.0% (100 basis points)	(44,046)	(21,224)
-0.5% (50 basis points)	22,023	10,612

The movements in profit are due to possible higher or lower interest costs from variable rate debt and cash balances.

Notes

Continued

17. FINANCIAL RISK MANAGEMENT OBJECTIVES (continued)

At balance date, the Group had the following mix of financial assets and liabilities exposed to variable interest rate risk:

(a) Cash flow interest rate risk (continued)

Consolidated Year ended	Note	Weighted average interest rate %	Non interest bearing	Floating interest rate	Floating interest maturing in			Total
					1 year or less	1 to 5 years	More than 5 years	
31 December 2010								
FINANCIAL ASSETS								
	7	4.70%	-	1,541,644	-	-	-	1,541,644
	8	n/a	4,295,487	-	-	-	-	4,295,487
Total financial assets			4,295,487	1,541,644	-	-	-	5,837,131
FINANCIAL LIABILITIES								
	13	n/a	1,082,492	-	-	-	-	1,082,492
	14	16.76%	-	-	2,400	24,453	-	26,853
	14	7.01%	-	-	6,075,900	-	-	6,075,900
	15	n/a	526,993	-	-	-	-	526,993
Total financial liabilities			1,609,485	-	6,078,300	24,453	-	7,712,238
Net exposure			2,686,002	1,541,644	(6,078,300)	(24,453)	-	(1,875,107)

Consolidated Year ended	Note	Weighted average interest rate %	Non interest bearing	Floating interest rate	Floating interest maturing in			Total
					1 year or less	1 to 5 years	More than 5 years	
31 December 2009								
FINANCIAL ASSETS								
	7	4.00%	-	4,612,205	-	-	-	4,612,205
	8	n/a	4,281,369	-	-	-	-	4,281,369
Total financial assets			4,281,369	4,612,205	-	-	-	8,893,574
FINANCIAL LIABILITIES								
	13	n/a	1,900,153	-	-	-	-	1,900,153
	14	5.83%	-	-	1,757,350	975,900	-	2,733,250
	15	n/a	462,915	-	-	-	-	462,915
Total financial liabilities			2,363,068	-	1,757,350	975,900	-	5,096,318
Net exposure			1,918,301	4,612,205	(1,757,350)	(975,900)	-	3,797,256



17. FINANCIAL RISK MANAGEMENT OBJECTIVES (continued)

(b) Credit risk

Credit risk arises from the financial assets of the Group, which comprise cash and cash equivalents and trade and other receivables. The Group's exposure to credit risk arises from potential default of the counter party, with a maximum exposure equal to the carrying amount of these instruments. Exposure at balance date is addressed in each applicable note.

The Group does not hold any credit derivatives to offset its credit exposure.

The Group trades only with recognised, creditworthy third parties and as such collateral is not requested nor is it the Group's policy to scrutinise its trade and other receivables. It is the Group's policy that all customers who wish to trade on credit terms are subject to credit verification procedures such as reviewing their industry reputation, financial position and credit rating. In addition, receivable balances are monitored on an ongoing basis with the result that the Group's exposure to bad debts is constantly managed.

There are no significant unprovided concentrations of credit risk within the Group.

(c) Liquidity risk

The Group's objective is to maintain a balance between continuity of funding and flexibility through the use of bank overdrafts and bank loans.

The Group's policy is to monitor the maturity of borrowings at all times. At 31 December 2010, 100% of the Group's debt will mature in less than one year (2009: 64%) however the majority of these facilities are in the process of renewal to extend the maturity dates.

Refer to the table below the heading 17 (a) Cash flow interest rate risk which reflects all contractually fixed pay-offs for settlement of financial liabilities and collection of financial assets. Trade payables and other financial liabilities generally originate from the financing of assets used in our ongoing operations such as investments in working capital eg inventories and trade receivables and investment in property plant and equipment. These assets are considered in the Group's overall liquidity risk. To monitor existing financial assets and liabilities as well as to enable an effective controlling of future risks, Cyclopharm monitors its Group's expected settlement of financial assets and liabilities on an ongoing basis.

The Group monitors rolling forecast of liquidity reserves on the basis of expected cash flow. At balance date the Group has no unused credit facilities (2009: \$3,716,750).

Consolidated Year ended	Note	Less than 6 months	6 months to 1 year	1 year to 5 years	Greater than 5 years	Total
31 December 2010						
Trade payables, third parties	13	1,082,492	-	-	-	1,082,492
Leases, third party	14	-	2,400	24,453	-	26,853
Secured bank loans, third party	14	-	6,075,900	-	-	6,075,900
31 December 2009						
Trade payables, third parties	13	1,900,153	-	-	-	1,900,153
Secured bank loans, third party	14	-	1,757,350	975,000	-	2,732,350

(d) Commodity price risk

The Group's exposure to commodity price risk is minimal.



17. FINANCIAL RISK MANAGEMENT OBJECTIVES (continued)

(e) Foreign currency risk

As a result of significant investment operations in Europe, the Group's Statement of Financial Position can be affected significantly by movements in the EURO / A\$ exchange rates. The Group does not hedge this exposure.

The Group also has transactional currency exposures. Such exposure arises from sales or purchases by an operating unit in currencies other than the unit's functional currency. Approximately 80% (2009: 85%) of the Group's sales are denominated in currencies other than the functional currency of the operating unit making the sale, whilst approximately 69% (2009: 51%) of costs are denominated in the unit's functional currency.

At 31 December 2010, the Group had the following financial instrument exposure to foreign currency fluctuations:

	Consolidated	
	2010	2009
	\$	\$
United States dollars		
Amounts payable	79,456	25,064
Amounts receivable	115,787	103,098
Euros		
Amounts payable	317,164	223,259
Amounts receivable	2,695,532	2,881,680
Canadian dollars		
Amounts payable	4,729	4,457
Amounts receivable	356,688	351,148
Net exposure	(2,766,658)	(3,083,146)

Management believe the balance date risk exposures are representative of the risk exposure inherent in the financial instruments.

Fair values

All of the Group's financial instruments recognised in the Statement of Financial Position have been assessed as at fair values.



17. FINANCIAL RISK MANAGEMENT OBJECTIVES (continued)

(e) Foreign currency risk (continued)

Foreign currency sensitivity

Currency risk is measured using sensitivity analysis. A portion of Cyclopharm's receivables and payables are exposed to movements in the values of those currencies relative to the Australian dollar. Cyclopharm management have determined that it is not cost effective to hedge against foreign currency fluctuations.

Cyclopharm is exposed to US Dollars (USD) and European Euro (Euro) movements. The following table details Cyclopharm's sensitivity to a 10% change in the Australian dollar against respective currencies with all other variables held constant as at reporting date for unhedged foreign exposure risk. A positive number indicates an increase in net profit/equity.

A sensitivity has been selected as this is considered reasonable given the current level of exchange rates and the volatility observed on a historic basis and market expectation for future movement.

	Consolidated	
	Increase in AUD of 10%	Decrease in AUD of 10%
	\$	\$
Euro		
31 December 2010		
Net profit	(162,471)	198,574
Equity increase/(decrease)	(162,471)	198,574
31 December 2009		
Net profit	(509,351)	622,538
Equity increase/(decrease)	(509,351)	622,538



18. COMMITMENTS

(a) Operating lease commitments

Future minimum rentals payable under non-cancellable operating leases are as follows:

	Consolidated	
	2010	2009
	\$	\$
Operating Lease Commitments		
Minimum lease payments		
Due not later than one year	432,441	170,266
Due later than 1 year & not later than 5 years	1,707,918	649,356
More than 5 years	1,332,720	-
Total operating lease commitments	3,473,079	819,622
Operating lease expenses recognised as an expense during the period:	165,897	148,579

- The Group has entered into commercial leases on office space within certain buildings. These leases have an average life of between 3 to 5 years with renewal options included in the contracts. There are no restrictions placed upon the lessee by entering into these leases.
- Cyclopet Pty Ltd has entered into a commercial lease for the PET Facility at Macquarie University Hospital. The lease has a term of 10 years and commenced upon commissioning of the Hospital in June 2010.
- The Group also has entered into commercial leases on motor vehicles that have an average life of approximately 3 to 5 years.

(b) Finance lease commitments

	Notes	Consolidated	
		2010	2009
		\$	\$
Finance Lease Commitments			
Minimum lease payments			
Due not later than one year	(i)	2,400	-
Due later than 1 year & not later than 5 years	(i)	24,453	-
More than 5 years		-	-
Total finance lease commitments		26,853	-

- (i) The Group also has entered into a commercial lease on motor vehicles that have a life of 5 years.

(c) Other commitments

	Notes	Consolidated	
		2010 \$	2009 \$
The company has the following other commitments:			
Not later than one year	(i) & (ii)	6,075,900	1,757,350
Due later than 1 year & not later than 5 years	-	-	975,900
More than 5 years	-	-	-
Total		6,075,900	2,733,250

- (i) Cyclopharm has a 15 month multi-option facility (MOF) provided by the National Australia Bank for \$5.1 million which expires in March 2011. At balance date \$5.1 million had been drawn down against this facility.
- (ii) Cyclopharm has an amortising bank bill facility provided by the National Australia Bank of \$1.35 million. At balance date \$975,900 was drawn down against this facility. Repayments under the amortising facility commenced in July 2010 and the entirety of the facility must be repaid by 31 July 2011.



18. COMMITMENTS (continued)

(d) Capital commitments

	Consolidated	
	2010	2009
	\$	\$
The company has the following capital expenditure commitments contracted for property, plant and equipment:		
Due later than 1 year & not later than 5 years	-	4,352,503
Total	-	4,352,503

19. RELATED PARTY DISCLOSURES

The consolidated financial statements include the financial statements of Cyclopharm and the subsidiaries as stated under the controlled entities note.

The following table provides the total amount of transactions that were entered into with related parties for the relevant financial year (for information regarding outstanding balances at year-end, refer to Note 8 Trade and other receivables, Note 13 Trade and other payables and Note 14 Interest bearing loans and borrowings):

CONSOLIDATED		Sales to related parties \$	Purchases from related parties \$	Transactions with related parties \$	Amounts owed by related parties \$	Amounts owed to related parties \$
CVC Venture Managers Pty Ltd	2010	-	-	16,189	-	2,963
	2009	-	-	39,160	-	1,870
VA Consulting Pty Ltd	2010	-	-	15,450	-	-
	2009	-	-	75,688	-	8,250
Macquarie Medical Imaging	2010	21,842	-	-	2,508	-
	2009	-	-	-	-	-

Ultimate parent entity

Cyclopharm Limited is the ultimate parent entity in the wholly owned group.

Terms and conditions of transactions with related parties

- During the year payments of \$16,189 (2009: \$39,160) were made to CVC Venture Managers (an entity of which Mr Sharman and Mr Gould are Non-Executive Directors) in relation to the rental of office space. Mr Gould does not receive any benefits from CVC Venture Managers.
- During the year payments of \$15,450 (2009: \$75,688) were made to VA Consulting Pty Ltd (an entity controlled by Mr Sharman). All payments related to Mr Sharman's role as a non-executive director.
- Cyclomedica Australia manufactures products that are sold to its overseas subsidiaries.
- Cyclopet Pty Ltd, a wholly owned subsidiary of Cyclopharm has a 20% interest in Macquarie Medical Imaging. Cyclopet manufactures products that are sold to Macquarie Medical Imaging.

19. RELATED PARTY DISCLOSURES (continued)

Controlled Entities

Name	Note	Country of Incorporation	Percentage of equity interest held	
			2009	2008
Cyclopharm Limited	1,2	Australia		
Controlled entities				
CycloPET Pty Ltd	2	Australia	100%	100%
Cyclomedica Australia Pty Limited	2	Australia	100%	100%
Cyclomedica Ireland Limited	3	Ireland	100%	100%
Cyclomedica Europe Limited	3	Ireland	100%	100%
Cyclomedica Germany GmbH	5	Germany	100%	100%
Cyclomedica Canada Limited	4	Canada	100%	100%
Allrad No 28. Pty Ltd	2	Australia	100%	100%
Allrad No 29. Pty Ltd	2	Australia	100%	100%

Notes

1. Cyclopharm Limited is the ultimate parent entity in the wholly owned group.
2. Audited by Russell Bedford NSW, Australia.
3. Audited by Moore Stephens Nathans, Republic of Ireland.
4. Audited by Schwartz Levitsky & Feldman & LLP, Toronto, Canada.
5. Audited by Bilzanzia GmbH Wirtschaftsprüfungsgesellschaft, Germany



20. EVENTS AFTER THE BALANCE DATE

There have not been any subsequent events after balance sheet date affecting the operating results of financial position of the Consolidated Group.

21. AUDITORS' REMUNERATION

The following total remuneration was received, or is due and receivable, by auditors of the Company in respect of:

	Consolidated	
	2010	2009
	\$	\$
Amounts received or due and receivable by Russell Bedford NSW and associated entities for:		
Audit and review of the financial statements	89,181	93,000
Other services:		
- tax compliance	11,030	12,730
- share registry	12,077	12,781
	112,288	118,511
Amounts received or due and receivable by auditors other than Russell Bedford NSW for:		
Audit of the financial statements	41,539	40,303
Other services	21,810	53,103
	63,349	93,406
Total auditors' remuneration	175,637	211,917

22. DIRECTOR AND KEY MANAGEMENT PERSONNEL DISCLOSURE

In accordance with the Corporations Amendment Regulations 2005 (No.4), the Company has transferred the remuneration disclosures required by *AASB 124: Related Party Disclosures* from the notes to the financial statements, to the Directors' Report under the heading of 'Remuneration Report'.



23. SHARE BASED PAYMENT PLANS

(a) Recognised share-based payment expenses

The expense recognised for employee services received in relation to share based payments during the year is shown in the table below:

	Consolidated	
	2010 \$	2009 \$
Expense arising from equity-settled share-based payment transactions (note 4)	96,177	74,283

The accumulated share based payment expense to 31 December 2010 was \$314,149 (2009: \$217,972).

(b) Type of share based payment plans

The share-based payment plan is described below. There have not been any modifications to the Long Term Incentive Plan ("Plan") following its approval by members at the Annual General Meeting held on 8 May 2007.

Shares

Long Term Incentive Plan ("Plan") Shares ("Shares") are granted to certain executive Directors and certain employees.

In valuing transactions settled by way of issue of shares, performance conditions and market conditions linked to the price of the shares of Cyclopharm Limited are taken into account. All shares issued have market performance conditions so as to align shareholder return and reward for the Company's selected management and staff ("Participants").

The Shares vest upon the satisfaction of certain performance conditions ("Hurdles") within the term ("Term") specified for Participants in the Plan. The Board has residual discretion to accelerate vesting (i.e. Reduce or waive the Hurdles) and exercise of Shares in the event of a takeover or merger or any other circumstance in accordance with the terms of the Plan.

Shares in relation to which Hurdles have not been satisfied (i.e. that do not vest) will lapse and will not be able to be exercised, except in the circumstances described below. Shares which have not vested will lapse where a Participant ceases employment with Cyclopharm other than on retirement, redundancy, death or total and permanent disablement or unless as otherwise determined by the Board in its absolute discretion.

Where a Participant has ceased employment with Cyclopharm as a result of resignation, retirement, redundancy, death or total and permanent disablement prior to the end of a performance period only Shares that have vested may be retained by the Participant on a pro-rata basis. If an option holder ceases employment for any reasons mentioned above prior to the first anniversary of the grant date, the Participant forfeits all entitlement to Shares.

LTIP Shares issued

At the Annual General Meeting held on 8 May 2007, Shareholders approved the Company's Plan.



23. SHARE BASED PAYMENT PLANS (continued)

Options

AASB 2 Share based Payment requires that the benefit to an employee arising from an employee share scheme such as the Cyclopharm Long Term Incentive Plan be treated as an expense in which the benefit is gained. No benefit to the employee arises from the Plan Shares as a corresponding loan applies to the issued Shares (although not required to be accounted for in the Financial Statements) instead the employee benefit is deemed to be the implied option ("Implied Option") arising from the Plan.

The International Financial Reporting Council have determined that where employee shares are issued under a non-recourse loan payment plan, the loan assets and the increment to share capital should not be recognised at grant date but rather, the transactions be treated as share options. Consequently the value of the discount which has been determined using a binomial pricing model will be charged to the Statement of Comprehensive Income over the vesting period. Other increments to share capital will be recognized as the share loans are settled by the relevant employees.

(c) Summary of shares granted

The following table illustrates the number of movements in share options during the current year:

	Consolidated 2010 Number	Consolidated 2009 Number
Balance at the beginning of the year	2,400,000	3,500,000
Granted during the year	-	-
Exercised during the year	-	-
Lapsed during the year	(1,700,000)	(1,100,000)
Balance at the end of the year	700,000	2,400,000
Exercisable at the end of the year	-	-
Number of recipients	-	-
Exercise price	-	-
Weighted average price	-	-
Exercise period from	-	-
To	-	-
Expiration day	-	-

23. SHARE BASED PAYMENT PLANS (continued)

(d) Option pricing models

The following assumptions were used to derive a value for the Implied Options granted using the Black Scholes Option model as at the grant date, taking into account the terms and conditions upon which the Shares were granted:

Exercise price per option	\$0.35
Number of recipients	\$1.00
Number of options	\$700,000
Grant Date	3/06/2008
Dividend yield	-
Expected annual volatility	38%
Risk-free interest rate	7.25%
Expected life of implied option (years)	4 years
Fair value per option	\$0.046
Share price at grant date	\$0.210
Model used	Black Scholes

Expected volatility percentages used for the Option pricing calculations were determined using historic data over 12 months and were adjusted to reflect comparable companies in terms of industry and market capitalisation. The Implied Options arising from the Plan are not listed and as such do not have a market value.

24. PARENT ENTITY DISCLOSURE

	Consolidated	
	2010	2009
	\$	\$
Financial Position		
Assets		
Current Assets	591,924	3,094,392
Non-current Assets	18,094,243	12,075,174
Total Assets	18,686,167	15,169,566
Liabilities		
Current Liabilities	5,505,951	1,496,730
Non-current Liabilities	1,264,500	2,066,976
Total Liabilities	6,770,451	3,563,706
Net assets	11,915,716	11,605,860
Equity		
Contributed equity	11,289,438	11,289,438
Employee equity benefits reserve	314,149	217,972
Retained Profits / (Accumulated losses)	312,129	98,450
Total Equity	11,915,716	11,605,860
Financial Performance		
Loss for the year	(213,679)	(968,804)

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