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To cite this article: Mark William Matthews (2011) A current review of empirical procedures of remote sensing in inland and near-coastal transitional waters, International Journal of Remote Sensing, 32:21, 6855-6899, DOI: [10.1080/01431161.2010.512947](https://doi.org/10.1080/01431161.2010.512947)

To link to this article: <http://dx.doi.org/10.1080/01431161.2010.512947>



Published online: 11 Aug 2011.



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## Review

### A current review of empirical procedures of remote sensing in inland and near-coastal transitional waters

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(Received 23 March 2010; in final form 24 July 2010)

The empirical approach of remote sensing has a proven capability to provide timely and accurate information on inland and near-coastal transitional waters. This article gives a thorough review of empirical algorithms for quantitatively estimating a variety of parameters from space-borne, airborne and *in situ* remote sensors in inland and transitional waters, including chlorophyll-*a*, total suspended solids, Secchi disk depth ( $z_{SD}$ ), turbidity, absorption by coloured dissolved organic matter ( $a_{CDOM}$ ) and other parameters, for example, phycocyanin. Current remote-sensing instruments are also reviewed. The theoretical basis of the empirical algorithms is given using fundamental bio-optical theory of the inherent optical properties (IOPs). Bands, band ratios and band arithmetic algorithms that could be used to produce common biogeophysical products for inland/transitional waters are identified. The article discusses the potential role that empirical algorithms could play alongside more advanced model-based algorithms in the future of water remote sensing, especially for near real-time operational monitoring systems. The article aims to describe the current status of empirical remote sensing in inland and near-coastal transitional waters and provide a useful reference to workers. It does not cover ‘inversion’ algorithms.

#### 1. Introduction

The recent launch of many new satellite instruments, and advances in computer technologies, has greatly increased the range of successful water-related remote-sensing applications and improved real-time monitoring of water quality and the rapid detection of environmental threats such as eutrophication and harmful algal blooms (HABs) (Mertes 2002, Ritchie *et al.* 2003, Glasgow *et al.* 2004, Power *et al.* 2005). Remote sensing offers substantial advantages over traditional monitoring methods, mainly because of the synoptic coverage and temporal consistency of the data, and has the potential to provide crucial information on inland and near-coastal transitional waters in countries where conventional water-quality monitoring programmes are either lacking or unsatisfactory (Navalgund *et al.* 2007). The great number of recent publications on the remote detection of a variety of biogeophysical parameters in inland and near-coastal transitional waters is testament to the rapidly growing interest in the subject (table 1). These waters are often optically complex and commonly known as ‘Case 2’ (Morel and Prieur 1977), being a function of at least

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three optically active constituents (phytoplankton, coloured dissolved organic matter (CDOM) and *tripton*), which may vary independently of one another. Therefore, these waters have more demanding requirements for instrument spectral resolution and sensitivity, atmospheric correction accuracy and water constituent retrieval algorithms (IOCCG 2000). Parameters often derived quantitatively using empirical methods from remotely sensed data include phytoplankton pigments such as chlorophyll-*a* (Chl-*a*) (Gitelson *et al.* 2009), cyanobacterial pigment phycocyanin (PC) (Ruiz-Verdú *et al.* 2008), concentration of total suspended solids (TSS) (Doxaran *et al.* 2009), absorption by coloured dissolved organic matter ( $a_{\text{CDOM}}$ ) (Kutser *et al.* 2005a), Secchi disk depth ( $z_{\text{SD}}$ ) or water clarity (Olmanson *et al.* 2008), turbidity (Petus *et al.* 2010) and water temperature (Giardino *et al.* 2001), among others. The empirical approach is distinguished from bio-optical model-based approaches by directly relating the remote-sensed signal to the parameter of interest using statistical techniques. The great number of recent studies using this approach proves that empirical algorithms have the capacity to provide reliable information on inland and transitional waters. Table 1 provides a comprehensive (but not exhaustive) overview of recent studies using empirical methods for quantitatively estimating a variety of biogeophysical parameters from space-borne, airborne and *in situ* remote-sensing instruments in inland and transitional waters. Table 1 gives information from each study on the location, the remotely sensed data type(s), the atmospheric correction procedure(s) (if applicable), the estimated water-quality parameter(s), the range of the parameter(s), the statistical technique(s) used to derive the algorithm(s), the band(s)/band ratios/arithmetic independent variable(s), the coefficient of determination ( $r^2$ ), the root mean square error and the sample size ( $N$ ). In some instances, only one best-performing or key algorithm from a study is included. The table is arranged by instrument in the following order: studies using dedicated ‘ocean colour’ sensors, for example, Moderate Resolution Imaging Spectroradiometer (MODIS), Medium Resolution imaging Spectrometer (MERIS) or simulations of these; studies using high-resolution sensors, for example, Landsat or simulations of these; and studies using hyper-/multispectral satellite and airborne platforms and *in situ* radiometers. The aim of this article is to describe the current status of empirical remote sensing in inland and transitional waters and identify remotely sensed band(s), band ratios and band arithmetic variables suitable for detecting specific parameters using certain instruments and statistical approaches in different water types. These are identified for their potential to provide common information products on inland and transitional waters using an empirically based approach. Special emphasis is given to explain the basis of these algorithms in terms of the inherent optical properties (IOPs), the absorption and backscattering coefficients of the optically active water constituents, phytoplankton, CDOM, detritus and minerals. In doing so, a theoretical basis for the algorithms is established using simple bio-optical theory: this is crucial for the cross-applicability and generalizability of the algorithms. The article begins with an overview of remote-sensing instruments as applicable to inland and transitional waters. It then gives a detailed review of empirical algorithms arranged according to the various biogeophysical parameters for various instruments and an explanation of their success in terms of simple bio-optical theory. The review ends by recommending empirical algorithms for common use and by discussing the role such algorithms could play alongside more advanced model-based approaches in the future of water remote sensing. This article should provide a useful reference of empirical algorithms for workers for a variety of ecosystem and monitoring applications.

Table 1. Recent remote-sensing studies in inland and near-coastal waters using empirical techniques (see Appendix A for a description of acronyms and abbreviations).

Year	Country	Study area	Data type*	Atmos. corr.	WQPs†	Data range ‡	Statistical technique	Bands/algorithm §	r <sup>2</sup>	RMSE ¶ (‰)	N	Reference
<i>Studies using multispectral ocean-colour sensors (including simulations)</i>												
2010	France	Bay of Biscay	SL MODIS (250 m)	MODIS Level 2G	TSS	0.3–145.6	Poly(2nd)	B2	0.96	61	74	Petus <i>et al.</i> (2010)
2009	Malawi	Lake Malawi	MODIS	MODIS Level 2G	Turb Chl- <i>a</i>	0.01–188.2 0.1–0.4	Poly(2nd) LR	B2 <i>R</i> 443/ <i>R</i> 551	0.95 0.58	378 –	74 5	Chavula <i>et al.</i> (2009)
2009	France	Gironde Estuary	MODIS AQUA (250 m)	MODIS Level 2G	TSS	77–2182	LR	B2/ <i>B</i> 1	0.82	245	75	Doxaran <i>et al.</i> (2009)
2009	USA	Fremont State Lakes	SL, MERIS	NA	Chl- <i>a</i>	~2–50	NLR	[ <i>R</i> 670 <sup>–1</sup> – <i>R</i> 710 <sup>–1</sup> ] × <i>R</i> 750	0.93	5.1 mg m <sup>–3</sup>	77	Gitelson <i>et al.</i> (2009)
2009	Russia	Taganrog Bay, Azov Sea	MERIS	MERIS Level 2	Chl- <i>a</i>	0.63–65.51	LR	<i>R</i> 708/ <i>R</i> 665	0.97	3.65 mg m <sup>–3</sup>	18	Moses <i>et al.</i> (2009a)
2009	Russia	Azov Sea	MERIS	MERIS Level 2	Chl- <i>a</i>	0.63–65.51	LR	[ <i>R</i> 665 <sup>–1</sup> – <i>R</i> 708 <sup>–1</sup> ] × <i>R</i> 753	0.95	5.02 mg m <sup>–3</sup>	18	Moses <i>et al.</i> (2009b)
2009	China	Dnieper Estuary Chaohu Lake	MODIS MODIS	MUMM 6S	Chl- <i>a</i> Chl- <i>a</i>	5.2–33.9	MLR	<i>R</i> 748/ <i>R</i> 667 LnB2 + LnB9– Ln(B3–B2)/B2	0.6 0.63	– –	7 40	Wu <i>et al.</i> (2009)
2008	Baltic Sea	Himmerfi-järden Bay	SL, MERIS	NA	z <sub>SD</sub>	0.25–1.2	MLR	LnB4/B10 + (B1 + B2)/B3 + B4	0.63	–	40	Kratzer <i>et al.</i> (2008)
2008	Germany	Lake Constance	MERIS	MIP	Chl- <i>a</i>	~3–6	LT-LR	<i>R</i> 490/ <i>R</i> 620	0.79	–	23	Odermatt <i>et al.</i> (2008)
2008	China	Poyang Lake	MODIS	MODIS Level 2G	TSS <i>a</i> <sub>CDOM</sub> z <sub>SD</sub>	0–10 0–0.3 0.06–2.84	Analytical MIP Analytical MIP LT-MLR	MIP B3, B1	– – 0.88	– – 37	– – 71	Wu <i>et al.</i> (2008)
2007	USA	Tampa Bay	LS TM MODIS	COST RTC	z <sub>SD</sub> Turb	0.32–2.16 0.9–8.0	LT-MLR LR	TMI, TM3 B1	0.83 0.73	20 –	25 43	Chen <i>et al.</i> (2007)

(Continued)

Table 1. (Continued.)

Year	Country	Study area	Data type*	Atmos. corr.	WQPs†	Data range ‡	Statistical technique	Bands/algorithm §	r <sup>2</sup>	RMSE ¶ (%)	N	Reference
2007	The Netherlands; 59 Lakes Spain		SI, MERIS	NA	PC	~0–1 100	SA	R709/R620	0.90	–	223	Simis <i>et al.</i> (2007)
2006	Gulf of Finland	Baltic Sea	MERIS	None	Chl- <i>a</i>	22–95	LR	L709/L665	0.87	22	51	Koponen <i>et al.</i> (2007)
2005	Italy	Lake Garda	MERIS	6S, DDV	TSS <i>a</i> <sub>CDOM</sub> Chl- <i>a</i>	2.9–12 1.29–2.15 ~2–11	LR LR LR	L709/(L560 + L665) L665/L490 L560/L665	0.92 0.96 0.49	16 5 1.20 mg m <sup>-3</sup>	51 51 7	Candiani <i>et al.</i> (2005)
2005	Italy	Lake Garda	SI, MIVIS MERIS	NA	Chl- <i>a</i>	3.5–8.9	LR	(L440–L780)/ (L480–L700)	0.69	9	22	Giardino <i>et al.</i> (2005)
2005	Netherlands	2 Lakes	MERIS SI, MERIS	None NA	Chl- <i>a</i> PC	0.2–2.5 0.8–79.8	LR SA	L489 R720/R620	0.83 0.94	26 19.7	31 33	Simis <i>et al.</i> (2005)
2004	Austria; Italy	4 Austrian Lakes; MERIS Lake Garda	MERIS	6S	Chl- <i>a</i>	1.0–5.6	LR	L665/L560	0.75	0.66 mg m <sup>-3</sup>	29	Floricioiu <i>et al.</i> (2004)
2004	Finland	Finnish Lakes	MODIS	None	Classes	1–4	Classifi- cation	B1	0.8 <sup>l</sup>	19.8% <sup>l</sup>	20391	Koponen <i>et al.</i> (2004)
2004	USA	Gulf of Mexico	MODIS	DOS	TSS	~1–55	LR	B1	0.89	2.18 mg l <sup>-1</sup>	52	Miller and McKee (2004)
2004	Italy	Lake Garda	MERIS	ELM	Chl- <i>a</i>	0.7–2.5	Poly(2nd)	L620/L709	0.76	0.27 mg m <sup>-3</sup>	12	Strömbeck <i>et al.</i> 2004
2003	USA	Gulf of Mexico	SI, SeaWiFS	NA	<i>a</i> <sub>CDOM</sub>	0.4–0.01	LT-LR	R443/R510	0.86	–	19	D'Sa and Miller (2003)
2003	Austria	3 Austrian Lakes	SI, ROSIS MERIS	6S	Chl- <i>a</i>	1.42–5.17	LR	L620/L490 + L560/L510	0.93	1.0 mg m <sup>-3</sup>	13	Floricioiu <i>et al.</i> (2003)
2003	Finland	Lake Hiidenvesi	SI, MERIS (AISA)	NA	Chl- <i>a</i>	6–44	LR	L705/L662	0.98	11.1	12	Kallio <i>et al.</i> (2003)
2003	The Netherlands	Lakes IJssel and Marken	SeaWiFS	SeaDAS	Chl- <i>a</i>	~5–160	Analytical 2 step	SeaWiFS	–	27	14	Vos <i>et al.</i> (2003)
2002	Sweden	Lake Märalen	SI, CASI MERIS	6S	TSS Chl- <i>a</i>	~10–250 2.5–18.9	Analytical 3 step SA	SeaWiFS R705/R664	– 0.88	38 –	14 570	Amnenberg <i>et al.</i> (2002)
					SPIM <i>a</i> <sub>CDOM</sub>	0.5–2.3 1.13–2.07	SA SA	R705/R664 R664/R550	0.83 –	0.34 mg m <sup>-3</sup> –	570 2	

(Continued)

Table 1. (Continued.)

Year	Country	Study area	Data type*	Atmos. corr.	WQPs†	Data range ‡	Statistical technique	Bands/algorithm§	r <sup>2</sup>	RMSE ¶ (‰)	N	Reference
2002	The Netherlands	IJssel Lagoon	SI, Spec. MERIS	NA	Chl- <i>a</i>	3–185	LR	L708/L664	0.96	8.3 mg m <sup>-3</sup>	114	Gons <i>et al.</i> (2002, 2005)
2002	Finland	Finnish Lakes	SI, MERIS (AISA)	None	Chl- <i>a</i>	1.3–100	LR	(L700–L781)/(L662–L781)	0.94	–	80	Koponen <i>et al.</i> (2002)
					<i>z</i> <sub>SD</sub>	0.4–7.0	LR	(L521–L781)/(L700–L781)	0.93	–	102	
					Turb	0.4–26 FNU	LR	L714	0.85	–	99	
2001	Sweden	Lakes Erken and Märalen	SI, CASI MERIS	6S	Chl- <i>a</i>	2.9–50.6	LR	R550	0.94	–	13	Flink <i>et al.</i> (2001)
					Chl- <i>a</i>	2.9–50.7	LR	R708/R678	0.84	–	13	
					Chl- <i>a</i>	2.9–50.8	MLR	R708/R678 + R643/R628	0.87	–	13	
					Chl- <i>a</i>	2.9–50.9	PCA; MLR	PC1 + PC2 + PC3 + PC4	0.96	–	13	
2001	Finland	Lakes	SI, MERIS (AISA)	None	Chl- <i>a</i>	1.3–100	LR	(L705–L754)/(L665–L754)	0.9	37	85	Härmä <i>et al.</i> (2001)
					TSS	0.7–23	LR	L705–L754	0.81	34	67	
					<i>z</i> <sub>SD</sub>	0.4–7.0	LR	(L490–L754)/(L620–L754)	0.83	35	85	
2001	Finland	11 Lakes	SI, MERIS (AISA)	MODTRAN	Chl- <i>a</i>	1–100	LR	L702/L674	0.91	29	88	Kallio <i>et al.</i> (2001)
					TSS	0.7–32	LR	R710	0.85	32	74	
					<i>z</i> <sub>SD</sub>	0.4–7	LR	(L492–L751)/(L622–L751)	0.86	30	103	
					<i>a</i> <sub>CDOM</sub>	1.2–14	LR	(L571–L607)/L607	0.84	20	47	
					Turb	0.4–26 FNU	LR	R710	0.93	23	105	
<i>Studies using multi-spectral high spatial resolution sensors</i>												
2009	China	Lake Chagan, Xinmiao, Kuli	LS TM	6S	<i>z</i> <sub>SD</sub>	0.22–0.79	LT-LR	TM3/TM2	0.91	0.03 m	15	Duan <i>et al.</i> (2009)
2009	Turkey	Lake Beysehir	ASTER	None	Chl- <i>a</i>	0.37–4.11	MLR	B1, B2, B3, B4	0.86	–	23	Nas <i>et al.</i> (2009)
2009	Turkey	Golden Horn Inlet	IKONOS	Erdaş ATCOR <sub>2</sub>	Chl- <i>a</i>	~0.1–2.14	MLR	B1, B2, B3, B4	0.88	–	9	Ormeci <i>et al.</i> (2009)

(Continued)

Table 1. (Continued.)

Year	Country	Study area	Data type*	Atmos. corr.	WQPs†	Data range ‡	Statistical technique	Bands/algorithm§	r <sup>2</sup>	RMSE ¶ (%)	N	Reference
2009	Japan	Lake Kasumigaura	SI, LS	NA	Chl- <i>a</i>	0–127	SDA	TM1, TM2, TM3	0.87	16.2 mg m <sup>-3</sup>	55	Oyama <i>et al.</i> (2009)
2008	Taiwan	Feitsui Reservoir	LS 7 ETM+	None	Chl- <i>a</i>	0.48–4.02	MLR	ETM1, ETM2, ETM3, ETM4, ETM5, ETM7	0.68	0.37 mg m <sup>-3</sup>	24	Chen <i>et al.</i> (2008)
					Chl- <i>a</i>	0.48–4.03	GEGA	ETM1, ETM4, ETM5, ETM7	0.79	0.30 mg m <sup>-3</sup>	24	
2008	Finland	Southern Lakes	LS ETM+	SMAC	zSD	0.5–5.5	LR	TM1/TM3	0.78	26.9	131	Kallio <i>et al.</i> (2008)
					acDOM Turb	1.0–12.2 0.6–15	NLR LR	TM2/TM3 TM3	0.83 0.86	22.3 28.7	29 80	
2008	USA	Minnesota's Lakes	LS TM, ETM+None	None	zSD	0.15–14.6	LT-MLR	TM1/TM3, TM1	0.71–0.96	14.1–40.6	13–278	Olmanson <i>et al.</i> (2008)
2008	Slovakia	Danube River	LS 7 ETM+	None	TSS	19.5–57.5	LR	ETM4	0.93	1.79 mg l <sup>-1</sup>	10	Onderka and Pekarova (2008)
2007	Turkey	Ömerli Dam	LS 7 ETM+	DOS	Chl- <i>a</i>	1.2–2.5	MLR	ETM1, ETM2, ETM3, ETM4	0.58	49	6	Alparslan <i>et al.</i> (2007)
					TSS	0.4–2.9	MLR	ETM1, ETM2, ETM3, ETM5	0.99	1	6	
					zSD	2.5–3.4	MLR	ETM1, ETM2, ETM3, ETM6	0.99	2	6	
2007	China	Lake Chagan	LS TM	None	Chl- <i>a</i>	6.3–58.2	LR	TM4/TM3	0.67	2.06 mg m <sup>-3</sup>	20	Duan <i>et al.</i> (2007)
2007	US	Charles River	Spec. IKONOS	NA None	Chl- <i>a</i> Turb	6.3–58.2 1.9–7.3	LT-LR LR	Ln(R700/R670) DN <sub>632-698</sub>	0.75 0.7	– –	54 3084	Hellweiger <i>et al.</i> (2007)
2006	France	Gironde Estuary	SI, SPOT	6S, DOS	TSS	10–2000	NLR	XS3/XS1	0.89	–	132	Doxaran <i>et al.</i> (2006)
2006	USA	Beaver Reservoir	SI, LS ETM+ LS TM	6S, DOS None	TSS Chl- <i>a</i>	10–2000 ~1.7–10	NLR ANN; LR	ETM4/ETM2 TM1, TM2	0.88 0.54	– 12.34 mg m <sup>-3</sup>	132 ~25	Sudheer <i>et al.</i> (2006)
					TSS	~0–11.5	ANN; LR	TM1, TM2, TM3, TM4	0.98	2.02 mg m <sup>-3</sup>	~25	

(Continued)

Table 1. (Continued.)

Year	Country	Study area	Data type*	Atmos. corr.	WQPs†	Data range ‡	Statistical technique	Bands/algorithm§	r²	RMSE ¶ (%)	N	Reference
2006	Central Europe	Lake Balaton	LS TM; ETM+DOS		Chl- <i>a</i>	~5–115	LMM	TM1, TM2, TM3 OR ETM1, ETM2, ETM3	0.95	–	11	Tyler <i>et al.</i> (2006)
2006	USA	Reelfoot Lake	LS 5 TM	Radiometric	TSS	~5–50	LR	TM3	0.89	–	11	Wang <i>et al.</i> (2006)
					Chl- <i>a</i>	66–189	MLR	TM2, TM3	0.71	–	18	
2005	USA	15 Minnesota lakes	LS TM	None	TSS	11.5–33.5	MLR	TM2, TM3, TM4	0.52	–	18	Brezonik <i>et al.</i> (2005)
					z <sub>SD</sub>	16–33	MLR	TM2, TM3	0.59	–	18	
					Turb	20.0–4.1	MLR	TM2, TM3	0.54	–	18	
					Chl- <i>a</i>	2.1–279	LT-MLR	TM1, TM1/TM3	0.88	–	15	
2005	USA	Pensacola Bay	LS 7 ETM+	Radiometric	Turb	0.3–155	LT-MLR	TM3	0.84	–	15	Han and Jordan (2005)
					z <sub>SD</sub>	0.15–4.4	LT-MLR	TM1, TM1/TM3	0.91	–	39	
					a <sub>CDOM</sub>	0.6–19.4	LT-MLR	TM1, TM1/TM4	0.77	–	15	
					Chl- <i>a</i>	1.1–23.2	LT-LR	Log ETM1/log ETM3	0.67	19	16	
2005	Finland; Sweden	Many lakes	ALI	ELM	a <sub>CDOM</sub>	0.68–11.13	NLR	L <sub>525-605</sub> /L <sub>630-690</sub>	0.73	–	30	Kutser <i>et al.</i> (2005)
2005	Central Europe	Lake Balaton	SI LS	NA	TSS	2–40.5	LR	TM2/TM3	0.88	–	10	Sváb <i>et al.</i> (2005)
2004	Taiwan	Techi Reservoir	LS TM	DOS	Cell density	~50–2400	LT-MLSR	TM1, TM2, TM3, TM4	0.73	–	120	Chang <i>et al.</i> 2004
2004	USA	New York Harbour	LS TM	Radiometric	z <sub>SD</sub>	~0.45–2	LT-LR	Log TM3	0.85	–	21	Hellweger <i>et al.</i> (2004)
2004	USA	Lake Erie	LS 5 TM	DOS	Chl- <i>a</i>	~5–50	LT-LR	Log(TM2/TM3)	0.78	–	16	Vincent <i>et al.</i> (2004)
					PC	~7.5–19	MLR	TM1, TM2, TM3, TM4; TM5	0.63	–	20	
2002	France	Gironde, Loire Estuaries	SI SPOT	NA	PC	~0.9–4.9	MLR	ETM1, ETM2, ETM3, ETM4, ETM5	0.78	15	30	Doxaran <i>et al.</i> (2003)
					Turb	~14.2–1.3	MLR	ETM3/ETM2	0.85	9	30	
					TSS	15–2500	NLR	XS3/XS1	0.93	–	200	
					TSS	15–2500	NLR	TM4/TM2	0.88	–	200	
			SI SeaWiFS	NA	TSS	15–2500	NLR	R865/R555	0.90	–	200	

(Continued)



Table 1. (Continued.)

Year	Country	Study area	Data type*	Atmos. corr.	WQPs†	Data range ‡	Statistical technique	Bands/algorithm§	r <sup>2</sup>	RMSE ¶ (%)	N	Reference
2002	France	Gironde Estuary	SI, SPOT	6S	TSS	35–2072	LR	XS3/XS1	0.93	<38	42	Doxaran <i>et al.</i> (2002b)
2002	The Netherlands	Southern Frisian lakes	SPOT HRV; L5 TM	L5MODTRAN-3	TSS	3–411	Analytical	(TM2 + TM3)/2 OR (XS1 + XS2)/2	0.99	–	–	Dekker <i>et al.</i> (2002)
2002	USA	Minnesota lakes	LS TM	ELM	zSD	~0.5–5	LT-MLR	TM1/TM3, TM1	0.71–0.92	~28	20–50	Kloiber <i>et al.</i> (2002)
2002	Finland	Gulf of Finland	LS TM, SAR	None	Chl- <i>a</i>	2.0–7.7	NN	TM1, ..., TM7, SAR	0.92	11.2	53	Zhang <i>et al.</i> (2002)
2001	Italy	Lake Guardia	LS TM	RTC – DOS	TSS	1.6–11.0	NN	TM1, ..., TM7, SAR	0.91	15.2	53	Brivio <i>et al.</i> (2001)
					zSD	0.67–4.2	NN	TM1, ..., TM7, SAR	0.95	7.3	53	
					Turb	1.0–7.5 FNUN	NN	TM1, ..., TM7, SAR	0.96	11.2	53	
					Chl- <i>a</i>	3.0–6.0	LR	(TM1–TM3)/TM2	0.82	37	5	
2001	Italy	Lake Iseo	LS TM	DOS	Chl- <i>a</i>	1.9–3.2	LT-MLR	L <sub>n</sub> TM1–L <sub>n</sub> TM2	0.68	49	5	Giardino <i>et al.</i> (2001)
					Chl- <i>a</i>	5.5–7.7	MLR	TM1, TM2	0.99	5.4	4	
2001	Southern Finland	Lakes	SI, LS (AISA)	NA	zSD	4.6–6.8	LR	TM1/TM2	0.85	45	4	Härmä <i>et al.</i> (2001)
					TSS	0.7–23	LT-LR	(TM1–TM4)/(TM3–TM4)	0.73	52	67	
					Turb	–	LT-LR	(TM1–TM4)/(TM3–TM4)	0.88	44	83	
2001	Sweden	Lake Erken	LS TM	6S	zSD	0.4–7.0	LR	(TM1–TM4)/(TM3–TM4)	0.81	34	85	Östlund <i>et al.</i> (2001)
					Chl- <i>a</i>	2.1–27.4	CHROM	TM1/TM1 + TM2 + TM3	0.93	–	19	
					Chl- <i>a</i>	2.1–27.4	LT-LR	Log(TM1/TM2)	0.88	–	19	
2001	China	Lake Taihu	LS TM	None	TSS	1.45–5.25	LR	TM1	0.95	–	19	Wang and Ma (2001)
					TSS	10–107	LT-MLR	L <sub>n</sub> (TM3 + TM4)/(TM1 + TM2)	–	–	15	
					zSD	0.2–0.5	PCA	L <sub>n</sub> TM1, L <sub>n</sub> TM2, L <sub>n</sub> TM3, L <sub>n</sub> TM5, L <sub>n</sub> TM7	–	–	15	
2000	Germany	Mecklenburg Lake District	LISS-III	MODTRAN	Chl- <i>a</i>	~2–63	LSU	LISS1, LISS2, LISS3	0.85	3.6 mg m <sup>-3</sup>	11	Thiemann and Kaufmann (2000)

(Continued)

Table 1. (Continued.)

Year	Country	Study area	Data type*	Atmos. corr.	WQPs†	Data range ‡	Statistical technique	Bands/algorithm§	r <sup>2</sup>	RMSE ¶(%)	N	Reference
1996	Israel	4 waste water reservoirs	SPOT	LOWTRAN 7	Chl- <i>a</i>	1.3–1600	BOM; PCA	XS1, XS2, XS3	–	–	4	Dor and Ben-Yosef (1996)
1996	Israel	Haifa Bay	Sl. LS	NA	TSS	2–195	BOM; PCA	XS1, XS2, XS3	–	–	4	
1995	Israel	Lake Kinneret	Sl. LS	NA	Chl- <i>a</i>	2–70	LT-LR	Log(TM3/TM1)	0.74	–	18	Gitelson <i>et al.</i> (1996)
1995	Israel	Lake Kinneret	Spec.	Fraser <i>et al.</i> (1992)	Chl- <i>a</i>	3.1–7.3	NLR	(TM1–TM3)/TM2	0.71	0.68 mg m <sup>–3</sup>	20	Mayo <i>et al.</i> (1995)
1995	Israel	Lake Kinneret	Spec.	NA	Chl- <i>a</i>	5.1–185	LR	Rmax/R670	0.95	3.4	41	Yacobi <i>et al.</i> (1995)
1989	USA	Lake Michigan	LS TM SPOT HRV	RTC Radiometric	Chl- <i>a</i> Chl- <i>a</i> z <sub>SD</sub>	5.1–186 5.1–187 0.6–2.0	LR LR LT-LR	RLH <sub>660-850</sub> TM4/TM3 XS3	0.96 0.79 0.83	3.2 4 20	41 40 11	Lathrop and Lillesand (1989)
1986	USA	Lake Michigan	LS TM	None	Turb TSS Chl- <i>a</i>	11.9–1.2 4.6–28.9 1.0–50.3	LT-LR LT-LR LT-LR	(XS2/XS1) + XS3 (XS2/XS1) + XS3 Ln TM2	0.88 0.93 0.98	29 19 1.04 mg m <sup>–3</sup>	11 11 13	Lathrop and Lillesand (1986)
<i>Studies using other hyperspectral, airborne and geostationary sensors</i>												
2009	Europe	North Sea	SEVIRI MSG	6S, SOS	TSS	0.5–9 12–0.54	LT-LR LT-LR	Ln TM2 Ln TM3	0.98 0.99	1.05 m 1.04	9 13	Neukermans <i>et al.</i> (2009)
2008	UK	Barton Broad	CASI-2	DOS	PC	<100 ~6–158	NLR LT-LR	VIS0.6 L710/L620	0.79	–	63	Hunter <i>et al.</i> (2008b)
2008	Spain, The Netherlands	64 freshwater lakes/ reservoirs	Spec.	NA	Chl- <i>a</i> PC	~4–63 ~0–640	LT-LR SA	R710/R670 R710/R620	0.96 0.92	18.3 47.5 mg m <sup>–3</sup>	13 352	Ruiz-Verdú <i>et al.</i> (2008)
					PC	~0–640	LR	0.5(R600 + R648)– R624	0.21	135.5 mg m <sup>–3</sup>	352	

(Continued)

Table 1. (Continued.)

Year	Country	Study area	Data type*	Atmos. corr.	WQPs†	Data range ‡	Statistical technique	Bands/algorithm§	r <sup>2</sup>	RMSE ¶(‰)	N	Reference
2007	Italy	Lake Garda	Hyperion	MODTRAN	PC Chl- <i>a</i>	~0–640 1.30–2.16	LR Analytical MIP	R650/R625	0.46 0.59	119.7 mg m <sup>-3</sup> 20	352 8	Giardino <i>et al.</i> (2007)
2006	China	Lake Tai	Spec.	NA	Turb Chl- <i>a</i>	0.95–2.13 20–190	Analytical MIP NLR	R719/R667	0.57 0.87	31 –	7 28	Jiao <i>et al.</i> (2006)
2006	USA	15 Minnesota Lakes	Spec.	NA	Chl- <i>a</i>	1.8–397	NLR	R700/R670	0.99	–	15	Menken <i>et al.</i> (2006)
2006	USA	Aquaculture ponds	Spec.	NA	Chl- <i>a</i>	107–3078	LR	(R740/R710)– (R740/R650)	0.78	319 mg l <sup>-3</sup>	~64	Zimba and Gitelson (2006)
2005	UK	Tamar Estuary	Spec.	NA	<i>a</i> <sub>CDOM</sub>	0.1–1.9	NLR	R400/R600	0.89	–	~43	Doxaran <i>et al.</i> 2005
2004	Finland	Gulf of Finland	Hyperion; ALIMODTRAN 4; ELM	Chl- <i>a</i>	1–1024	SAM; BOM	–	–	–	–	–	Kutser (2004)
2004	Germany	Lake Constance	CHRIS	RTM	Chl- <i>a</i>	1.4–4.4	Analytical MIP	–	–	–	17	Miksa <i>et al.</i> (2004)
2003	Australia	Moreton Bay	Hyperion	MODTRAN	TSS <i>a</i> <sub>CDOM</sub> Chl- <i>a</i>	1.1–2.6 0.13–0.30 1.0–19.9	Analytical MIP Analytical MIP Analytical MIP	– – –	– – –	– – –	17 17 –	Brando and Dekker (2003)
					0.13–0.75 3.4–46.3 13–985	Analytical MIP Analytical MIP Poly(3rd)	– – R850/R550	– – 0.97	– – –	– – 34		
2002	France	Gironde Estuary	Spec.	NA	TSS	~0.1–1.5	LR	R670/R412	0.99	–	8	Doxaran <i>et al.</i> (2002a)
2000	Scotland	Clyde Sea	Spec.	NA	<i>a</i> <sub>CDOM</sub>	3–185	SA	R704/R672	0.95	3 mg m <sup>-3</sup>	114	Bowers <i>et al.</i> (2000)
1999	The Netherlands	Lakes, rivers and estuaries	Spec.	NA	Chl- <i>a</i>	1–171	MLR	dR429 + dR695 + (dR429 × dR695)	0.5	–	19	Gons (1999)
1998	USA	Nebraska Sand Hills Lakes	Spec.	ELM	Chl- <i>a</i>	1–82	MLR	dR429 + dR628 + d695	0.69	–	30	Fraser (1998)

(Continued)

Table 1. (Continued.)

Year	Country	Study area	Data type*	Atmos. corr.	WQPs†	Data range‡	Statistical technique	Bands/algorithm§	r <sup>2</sup>	RMSE ¶(%)	N	Reference
1998	USA	Carter Lake	Spec.	NA	Chl- <i>a</i>	20–280	LR	RLH <sub>670-850</sub>	0.86	±5.3 mg m <sup>-3</sup>	35	Schalles <i>et al.</i> (1998)
1994	USA	Tennessee Reservoirs	AMMS	None	Chl- <i>a</i>	21–280	LR	SUM <sub>670-850</sub>	0.87	±5.3 mg m <sup>-4</sup>	36	Dierberg and Carriker (1994)
					Chl- <i>a</i>	2–79	LR	R700/R680	0.95	2.19 mg m <sup>-3</sup>	29	
1994	Israel	Lake Kinneret	Spec.	NA	Chl- <i>a</i>	2–79	LR	FLH <sub>663-700</sub>	0.85	2.76 mg m <sup>-3</sup>	29	Gitelson <i>et al.</i> (1994)
					Turb	1.1–11	LR	R700/R680	0.88	0.93 NTU	29	
					Chl- <i>a</i>	1–46	LR	L694/L679	0.84	2 mg m <sup>-3</sup>	34	
					Chl- <i>a</i>	1–46	LR	SLH <sub>665-752</sub>	0.86	1.9 mg m <sup>-3</sup>	34	
1993	Europe	> 20 Inland water bodies	Spec.	NA	Turb	1.1–10	LR	L694/L679	0.77	1.0 NTU	34	A. Gitelson <i>et al.</i> (1993)
					Chl- <i>a</i>	3.1–7.3	LR	SUM <sub>670-730</sub>	0.84	0.69 mg m <sup>-3</sup>	20	
					Chl- <i>a</i>	3.1–7.3	LR	FLH <sub>670-730</sub>	> 0.73	0.77 mg m <sup>-3</sup>	20	
1992	Germany	Laboratory, rivers and lakes	SI, LS MSS	NA	Chl- <i>a</i>	0.1–350	NLR	R700/R675	> 0.88	< 2 mg m <sup>-3</sup>	> 383	Mittenzwey <i>et al.</i> (1992)
					TSS	0.1–66	NLR	(R560–R520)/(R560 + R520)	0.86	1.79 mg l <sup>-1</sup>	66	
					<i>a</i> <sub>CDOM</sub>	0.1–12	NLR	((R480–R700/R675)–R520)/((R480 + R700)/R675) + R520)	0.9	0.25 mgC m <sup>-3</sup>	–	
					Chl- <i>a</i>	30–150	MLR	MSS6/MSS4 + MSS5 + MSS6	0.88	< 4.47 mg m <sup>-3</sup>	134	
					Chl- <i>a</i>	5–350	MLR	L705/L670	0.98	–	94	

Notes: Coefficients of determination and variables other than spectral bands have not been included. Only algorithms with highest performing correlation coefficients are presented.  
\*SI. = simulated. †WQPs = Water-quality parameters. ‡Units for Chl-*a*, PC = mg m<sup>-3</sup>; TSS, SPIM = g m<sup>-3</sup>; *a*<sub>CDOM</sub> = m<sup>-1</sup> (at 440 nm unless otherwise stated); *z*<sub>SD</sub> = m; Turb = NTU (nephelometric turbidity units), unless otherwise indicated. §*Lx* = radiance at wavelength *x*; *Rx* = reflectance at wavelength *x*; DN = digital number; satellite bands are prefixed by the abbreviated sensor's name. RMSE¶ = root mean square error in percentage unless otherwise indicated. †Total classification accuracy/error.

## 2. Remote-sensing instruments and their application to inland and near-coastal transitional waters

Passive remote-sensing instruments, whether handheld or mounted on aircraft or satellites, measuring the light in the visible and near-infrared (NIR) part of the electromagnetic spectrum (400–1000 nm) are most often used for water-related applications. The optically active water constituents, including phytoplankton, *tripton* made up of detritus and minerals, CDOM (also called *gelbstoff* or yellow substances) and water itself, all have an impact on the optical signature of water in the visible wavelengths. Viruses, bacteria, bubbles and other aqueous particles may also be significant (see Stramski *et al.* 2004) but are generally not routinely detected. The water-leaving radiance is modified through the backscattering ( $b_b$ ) and absorption ( $a$ ) of light by these constituents (called the IOPs) (Preisendorfer 1976). Absorption by phytoplankton,  $a_\phi$ , *gelbstoff* and detritus,  $a_{dg}$ , and water,  $a_w$ , are well defined (figure 1) and can be used to explain the causal relationships between the observed remote-sensing reflectance and the biogeophysical parameter(s) of interest in terms of bio-optical theory (see Kirk 1994). The backscattering coefficients for water,  $b_{bw}$ , minerals,  $b_{bm}$ , phytoplankton,  $b_{b\phi}$ , and particulate matter,  $b_{bp}$  (negligible for *gelbstoff* and detritus) may be used in the same way. Strong absorption by water at wavelengths  $> \sim 750$  nm (Buiteveld *et al.* 1994) effectively masks out the signals from other constituents except in highly turbid water where scattering by minerals overwhelms absorption by water. Therefore, wavelengths between 400 and 750 nm generally contain the most information on the water constituents, which is detectable by remote-sensing instruments, with the exception of highly turbid water where the signal in the NIR is also useful.

Portable field spectroradiometers typically used for *in situ* calibration/validation purposes provide multi- or hyperspectral measurements (up to 1 nm resolution) of the upwelling radiance above the water, or just beneath the surface, and the downwelling irradiance above the surface. There are a wide range of spectroradiometers

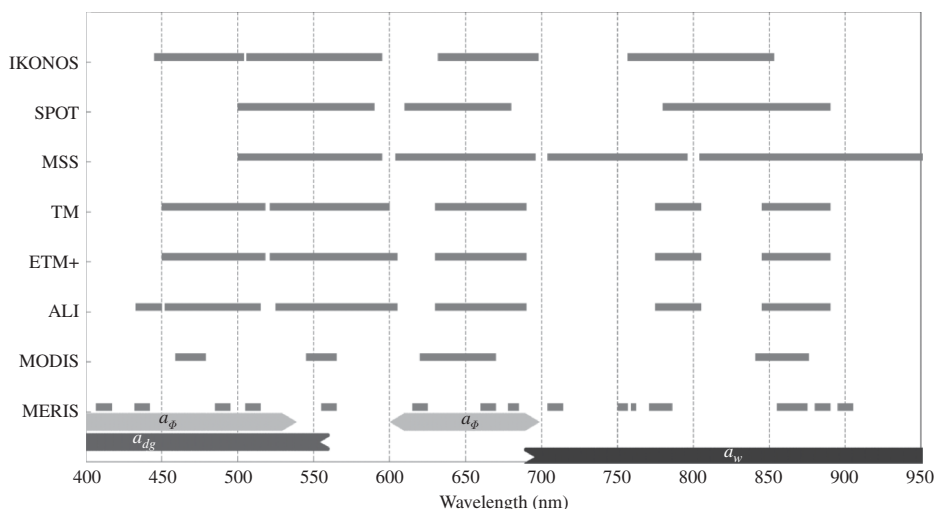


Figure 1. The spectral position of various satellite instruments in relation to the location of the maximum influence of absorption by phytoplankton,  $a_\phi$ , detritus and *gelbstoff*,  $a_{dg}$ , and water,  $a_w$ . The bands plotted for MODIS are the 250 and 500 m bands.

available. Commonly used instruments include the PR-650 SpectraColorimeter (Photo Research, Chatsworth, CA, USA) (Gons *et al.* 2002, Giardino *et al.* 2007, Simis *et al.* 2007); the TriOS-RAMESS radiometers (TriOS Optical Sensors, Germany) (Doxaran *et al.* 2005, Ruddick *et al.* 2006, Neukermans *et al.* 2009); and the ASD FieldSpec spectroradiometer (Analytical Spectral Devices, Inc. Boulder, CO, USA) (Ruiz-Verdú *et al.* 2005, Jiao *et al.* 2006, Simis *et al.* 2007). *In situ* hyperspectral reflectance/radiance measurements have been essential not only for calibration/validation of the water-leaving reflectance derived from satellite/airborne measurements, but also for the derivation of empirical algorithms (e.g. Gitelson *et al.* 1993, Doxaran *et al.* 2002a, Zimba and Gitelson 2006). The findings of these and similar studies form the basis for algorithm development and are therefore a vital component in the development of remote-sensing systems.

Space and airborne remote-sensing instruments used for inland and near coastal water-related applications must meet the minimum spectral, spatial, temporal, radiometric and signal-to-noise ratio (SNR) requirements. The detection of certain water constituents, such as the phytoplankton pigment Chl-*a*, necessitates that the remote-sensing reflectance be resolved in sufficient detail for the application of suitable detection algorithms. The ground resolution of a pixel for larger inland waters ( $> \sim 1 \text{ km}^2$ ) should be a few hundred metres or less, several times larger than the dimensions of the target. Also, the sampling frequency should be regular, especially when considering that cloud cover may substantially reduce the number of useful images. Higher radiometric resolutions and SNRs are also required to describe sufficiently the low range of reflectance values over water. In large, these sensor requirements are dictated by the intended application: ecosystem analysis applications generally require higher sampling frequencies to resolve system changes occurring over short timescales (days/weeks), and high SNRs to improve confidence limits; course change detection applications with larger signals, such as eutrophic status determination, are generally accommodated with more coarse radiometric resolutions and lower overpass frequencies. Therefore, the selection of the remote-sensing instrument is based on the desired application, for example, operational PC detection from currently available high spatial resolution satellite sensors may not be feasible (see section 4.5.3).

Airborne platforms typically carry multi- or hyperspectral spectrometers capable of capturing an almost continuous spectrum of the surface reflectance. Examples of airborne sensors commonly used in inland waters are the Airborne Imaging Spectrometer for Applications (Härmä *et al.* 2001, Kallio *et al.* 2003) and the Compact Airborne Spectrographic Imager (Ammenberg *et al.* 2002, Hunter *et al.* 2009). The data from these sensors are particularly useful, because the spectral widths and positions can normally be adjusted to suit the intended application or resampled to simulate multispectral satellite sensors that have broader and fewer bands. The very high spectral resolution presents opportunities for applications that are not feasible with sensors with few and broad bands, such as the determination of algal species composition through the detection of specific algal pigments, for example, PC pigment present in cyanobacteria (Richardson 1996, Gege 1998, Hunter *et al.* 2008a). The spatial resolution of most airborne sensors is also high, given the low altitude at which the images are acquired, and the contribution from the atmosphere is often presumed to be negligible, which means that atmospheric correction may sometimes be ignored. However, atmospheric correction is nevertheless required when using algorithms based on  $R_{rs}$ . Furthermore, airborne campaigns can be timed to coincide

with events such as blooms, tides, floods and other episodic events providing more flexibility than satellite sensors. There are numerous recent examples where airborne sensors have been used successfully for the detection of various water-quality parameters in lakes (e.g. Dierberg and Carriker 1994, Hakvoort *et al.* 2002, Koponen *et al.* 2002, Floricioiu *et al.* 2003, Hunter *et al.* 2009, table 1). However, the difficulty in obtaining temporally and spatially consistent data as acquired from satellites, and the relatively high cost of campaigns, makes this approach generally unsuited to frequent monitoring applications, especially in the developing world.

Satellite platforms offer substantial advantages over airborne platforms mainly because of the temporal and spatial consistency, larger area coverage and reduced data cost borne over a large user community (Rees 2001). There are a number of current and scheduled Earth-observation satellites that may be used for water-quality monitoring (see table 2). Hyperspectral satellite-based sensors are only now progressing beyond the experimental phase owing to constraints related to instrument design (SNR), difficulties with atmospheric correction and the large quantities of data that must be transmitted and stored. The first experimental hyperspectral sensor to orbit the Earth in space was the Hyperion imaging spectrometer, launched in 2001, which has been used for deriving detailed water-quality parameters in inland and near-coastal transitional waters (Brando and Dekker 2003, Kutser 2004, Giardino *et al.* 2007). Hyperion has 220 bands between 0.4 and 2.5  $\mu\text{m}$ , a spatial resolution of 30 m and an overpass time equivalent to the Landsat sensors (16 days). Hyperion is now near the end of its lifespan and does not acquire images routinely. The recently launched experimental Hyperspectral Imager for the Coastal Ocean (HICO) on board the International Space Station is designed specifically for monitoring the littoral coastal environment and has been acquiring imagery since 25 September 2009. HICO has a high SNR and a ground resolution of 90 m, making it suitable for lakes (Corson *et al.* 2008). Similar sensors are likely to be used increasingly in the future (e.g. the Enmap Hyperspectral Imager scheduled for launch in 2012). High- to medium-resolution spectrometers with fewer bands than hyperspectral sensors, such as the experimental Compact High Resolution Imaging Spectrometer (CHRIS) on board the Proba-1 platform, have also been used for water-quality monitoring (Miksa *et al.* 2004, Ruiz-Verdú *et al.* 2005). CHRIS has 19 bands in 'water mode' ideally positioned for water-related applications, a spatial resolution of 18 m (swath width is only about 14 km) and an image acquisition frequency of about 7 days. However, as CHRIS is only an experimental sensor, actual data acquisition may be infrequent due to conflicting demands of different sites for images, and there are also substantial challenges related to atmospheric correction. It is likely that other spectrometers with high spectral definition and spatial resolution, and fairly regular image acquisition, will be used increasingly in the future.

Multispectral sensors such as IKONOS, LISS 3 and 4, SPOT 4 and 5, Landsat 5 and 7 and ALI have few, broad bands and high spatial resolutions (4–30 m) and are primarily designed for terrestrial applications (see table 2 and figure 1). Nonetheless, there are a great number of successful studies that use these instruments to estimate a variety of water-quality parameters in inland/transitional waters (e.g. Thiemann and Kaufmann 2000, Dekker *et al.* 2001, Chang *et al.* 2004, Vincent *et al.* 2004, Wang *et al.* 2004, Kutser *et al.* 2005a, Doxaran *et al.* 2006, Hellweger *et al.* 2007, table 1). These instruments offer the advantage of being able to view even very small lakes with high spatial definition. However, the spectral and radiometric configuration (broad bands, low SNR) of these instruments generally limits their usefulness to course change-detection studies: the broad spectral bands impede the detection of certain

Table 2. Current Earth-observation satellite sensors, which may be used for water-quality assessments in inland and near-coastal waters showing resolution specifications and full names. Italics indicate scheduled sensors.

Satellite	Sensor	Spectral resolution ( $\mu\text{m}$ )	Spatial resolution	Temporal resolution	Example of study
LM900	IKONOS	0.45–0.85 (4 bands)	4 m	3/5 days	Hellweger <i>et al.</i> (2007)
IRS-P6	LISS 4	0.52–0.68 (3 bands)	5.8 m	5 days	–
SPOT 5	HRG	0.48–1.75 (5 bands)	10 m	26 days	Dekker <i>et al.</i> (2002)
Proba-1	CHRIS	0.415–1.050 (19 bands)	18 m	~7 days	Miksa <i>et al.</i> (2004)
SPOT 4	HRVIR	0.50–0.89 (3 bands)	20 m	26 days	Lathrop and Lillesand (1989)
IRS-P6	LISS 3	0.52–1.70 (4 bands)	23.5 m	24 days	Thiemann and Kaufmann (2000)
<i>EnMAP</i>	HIS	<i>0.420–2.450 (200 bands)</i>	<i>30 m</i>	<i>4 days</i>	–
EO-1	Hyperion	0.4–2.5 (220 bands)	30 m	16 days	Giardino <i>et al.</i> (2007)
EO-1	ALI	0.43–2.35 (9 bands)	30 m	16 days	Kutser <i>et al.</i> (2005)
Landsat 5	TM	0.45–2.35 (6 bands)	30 m	16 days	Vincent <i>et al.</i> (2004)
Landsat 7	ETM+	0.45–2.35 (8 bands)	30 m	16 days	Olmanson <i>et al.</i> (2008)
ISS	HICO	0.3–1.0 (128 bands)	100 m	–	–
Terra/Aqua	MODIS	0.620–0.876 (2 bands)	250 m	1–2 days	Chen <i>et al.</i> (2007)
EnviSAT	MERIS	0.412–0.900 (15 bands)	~300 m	2–3 days	Giardino <i>et al.</i> (2005)
<i>Sentinal 3</i>	<i>OLC</i>	<i>0.413–1.020 (16 bands)</i>	<i>~300 m</i>	<i>2–3 days</i>	–
IRS-P4	OCM	0.400–0.885 (8 bands)	360 m	2 days	–
<i>COMS</i>	<i>GOCI</i>	<i>0.400–0.900 (8 bands)</i>	<i>500 m</i>	<i>~15 min</i>	–
<i>TRAQ</i>	<i>OCAPI</i>	<i>0.320–2.13 (8 bands)</i>	<i>4 km</i>	<i>14 min</i>	–
SeaWiFS	WiFS	0.402–0.885 (8 bands)	1 km	2 days	Vos <i>et al.</i> (2003)

Notes: ALI, Advanced Land Imager; CHRIS, Compact High Resolution Imaging Spectrometer; COMS, Communication, Ocean, and Meteorological Satellite; EnMAP, Environmental Mapping and Analysis Program; ETM, Enhanced Thematic Mapper; GOCI, Geostationary Ocean Colour Imager; HICO, Hyperspectral Imager for the Coastal Ocean; HIS, Hyperspectral Imaging Spectroradiometer; HRVIR, High Resolution Visible and InfraRed imaging instrument; HRG, High Resolution Geometric imaging instrument; IKONOS, Derived from the Greek word for ‘image’; ISS, International Space Station; LISS, Linear Imaging Self-Scanning Sensor; MODIS, Moderate Resolution Imaging Spectrometer; MERIS, Medium Resolution Imaging Spectrometer; OCAPI, Optical Carbonaceous and anthropogenic Aerosols Pathfinder Instrument; OCM, Ocean Colour Monitor; OLC, Ocean and Land Colour Imager; SeaWiFS, Sea-viewing Wide Field-of-view Sensor; TM, Thematic Mapper; TRAQ, Tropospheric composition and Air Quality.

parameters, and the absence of NIR bands (in some sensors) makes atmospheric correction more challenging. The low acquisition frequencies (with the exception of constellations, e.g. SPOT) mean that they are better suited to event-scale rather than



frequent change-detection applications. In general, these instruments are used for deriving site-and-time parameter-specific empirical algorithms – although the cross-applicability of algorithms for certain parameters has been shown (e.g. Dekker *et al.* 2002, Doxaran *et al.* 2006, Olmanson *et al.* 2008). For operational purposes, data cost from these instruments is a significant concern – although there are exceptions such as Landsat, which is available free of charge including for countries in the developing world, although some of the data are unusable due to the SLC (scan line corrector) failure and the archive is sporadic (NASA 2008).

Typical ‘ocean-colour’ sensors more suited to real-time operational and detailed parameter retrieval, such as MODIS, SeaWiFS, OCM and MERIS, have high acquisition frequencies, bands ideally positioned for the detection of water constituents and atmospheric correction and high SNRs (see table 2 and figure 1). The obvious disadvantage of these sensors for inland waters is their lower spatial resolution, typically between 250 m and 1000 m in full-resolution mode. The spatial resolution of SeaWiFS and MODIS ocean-colour bands (1 km) is too low for all but the largest inland lakes (Vos *et al.* 2003, Chavula *et al.* 2009). MODIS’s two broad high-resolution bands (250 m) have been used successfully for monitoring TSS and for water classification in inland lakes and estuaries (Koponen *et al.* 2004, Miller and McKee 2004, Chen *et al.* 2007, Doxaran *et al.* 2009, Petus *et al.* 2010). However, its ability to detect other crucial parameters, such as phytoplankton pigments (Chl-*a*), is limited. MERIS is perhaps the best suited of these sensors for inland water-quality monitoring with a full resolution of  $\sim 260 \times 290$  m at nadir and 15 bands in the visible and NIR. There are several examples using MERIS in inland and near-coastal waters (e.g. Floricioiu *et al.* 2004, Giardino *et al.* 2005, Koponen *et al.* 2007, Kratzer *et al.* 2008, Odermatt *et al.* 2008, Moses *et al.* 2009a). The availability of a standard MERIS Level 2 product suited for use in coastal waters (Schiller and Doerffer 1999, 2005) and products for regional case 2 waters and lakes (Doerffer and Schiller 2008a, b) is very convenient for a variety of studies – although these algorithms require parameterization to work in different regions. Recent research shows that MERIS may even be used in small lakes with equal or more effectiveness than Landsat, which makes it arguably the current optimal sensor for inland water monitoring (Matthews *et al.* 2010). Data from these ocean-colour sensors are also available free for research purposes, making them a more viable option for scientists, particularly in the developing world.

The proposal of using geostationary satellite sensors to monitor parameters such as TSS has already been proved using the SEVIRI (spinning enhanced visible and infrared imager) sensor on board METEOSAT (meteorological satellite) Second Generation (Neukermans *et al.* 2009, Salama and Shen 2010). The first image from a dedicated ocean-colour geostationary sensor was acquired on 13 July 2010 by the Global Ocean Colour Imager (Korean Aerospace Research Institute) (Kang *et al.* 2006). Future sensors such as the Optical Carbonaceous and anthropogenic Aerosols Pathfinder Instrument (CNES/ESA) (Le Naour *et al.* 2006) and the Geostationary Coastal and Air Pollution Events sensors (NASA) (Al-Saadi *et al.* 2009) will be capable of providing almost continuous data ( $\sim 15$  min) – the main challenges are related to radiometric resolution and sensitivity due to the distance from the target (Kang *et al.* 2006).

### 3. Quantitative remote sensing of biogeophysical water-quality parameters

Broadly, there are two approaches for deriving water-quality products from remotely sensed data: the empirical and model-based approaches. The model-based (or analytical) approach seeks to model the remote-sensing reflectance,  $R_{rs}(0+)$  (or the

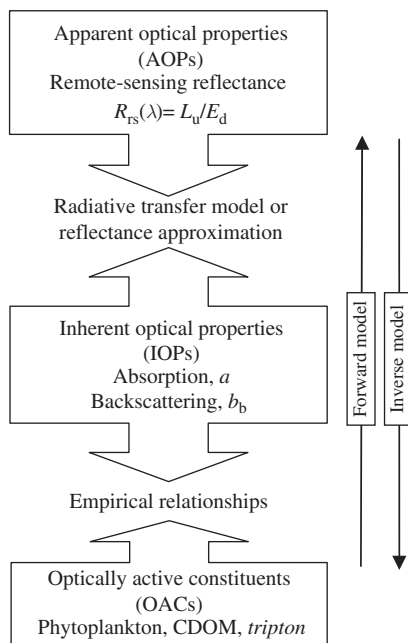


Figure 2. Simplified schematic diagram of the forward and inverse models used in model-based approaches.

reflectance at the top of the atmosphere,  $R_{TOA}$ ) in terms of the water IOPs through radiative transfer modelling (see Dekker *et al.* 2001 for an overview) (see figure 2).

The ‘forward’ model derives  $R_{rs}(0+)$  from the water IOPs using a bio-optical model and an approximation of the radiative transfer equation, called the reflectance approximation (Morel and Prieur 1977, Zaneveld 1995), or through direct solution of the RTE (Radiative Transfer Equation) using models such as Hydrolight (Mobley 1994). The  $R_{TOA}$  can then be modelled using radiative transfer calculations for the atmosphere through codes such as 6S (Vermote *et al.* 1997) or MODTRAN (Kneizys *et al.* 1988). The ‘inverse’ model solves the IOPs (or concentrations of optically active water constituents) from reflectance measured at the top of the atmosphere by satellites or from *in situ* measurements of  $R_{rs}(0+)$ . The ‘inversion’ problem may be solved using any of a variety of mathematical optimization or multiple non-linear regression procedures, such as artificial neural networks, to produce ‘analytical’ or ‘inversion’ algorithms (IOCCG 2000, 2006). This allows a number of parameters to be solved simultaneously, most often  $a_\Phi$  (which may be converted to the concentration of Chl-*a*), backscattering from suspended solids,  $b_{bp}$ , and  $a_{CDOM}$ , independent of simultaneously acquired experimental data (e.g. Odermatt *et al.* 2008). The European Space Agency’s MERIS Lakes and Case 2 Processors, such as the Eutrophic Lakes Processor, are examples of operational inversion-type algorithms for use in inland waters (Doerffer and Schiller 2008a, b). The main concerns with these kinds of algorithms are their sensitivity to errors from atmospheric correction procedures (e.g. Lee *et al.* 2002) and the existence of non-unique or ambiguous solutions arising from the additive nature of the IOPs and the consequences of using a ratio in the reflectance approximation (Defoin-Platel and Chami 2007). The analytical approach is generally complex and ideally requires measurements and knowledge of local/regional IOPs to develop a

robust forward model. This often requires substantial fieldwork and algorithm training and computing time. As this review focuses on the empirical approach, analytical algorithms are beyond the scope of this article and reviews can be found elsewhere (see IOCCG 2000, 2006). Closely related to the analytical approach is what is often called the semi-analytical approach, which uses algebraic solutions of the reflectance approximation to derive biogeophysical parameters (e.g. Gons 1999, Gons *et al.* 2002, Simis *et al.* 2005 and others). Some studies using the semi-analytical approach are included in this review because they have rather simple algebraic solutions and are also useful for understanding the causal relationship between the remote-sensing reflectance, the IOPs and the parameter of interest.

In contrast to the analytical approach, empirical algorithms are relatively simple to derive and use: simultaneously acquired experimental sets of limnological, atmospheric and remotely sensed data are used to derive normally site-and-time specific algorithms for a certain parameter using statistical regression techniques. These algorithms generally produce robust results for the areas and data sets from which they are derived. Figure 3 shows a simplified schematic diagram of the main steps used in the empirical approach.

The first step involves the collection of simultaneous sets of remotely sensed, atmospheric and biogeophysical water-quality parameter data (this may also include IOPs, e.g. the total absorption coefficient,  $a$ , or backscattering from particulate matter,  $b_{bp}$ ). The acquisition of all these data simultaneously requires substantial effort especially when considering that cloud cover may significantly reduce the number of successful matchups. After collection, the data must be processed to a form suitable for the intended application. This may include atmospherically correcting the top of the atmosphere radiances to surface reflectances, which may be performed using a variety of techniques (see table 1). The next step is to determine correlations between the remotely sensed data and the water-quality parameter of interest, which may be achieved using linear, multiple linear, non-linear or other statistical models. This leads to the derivation of an empirical algorithm, which may be used to estimate the spatial

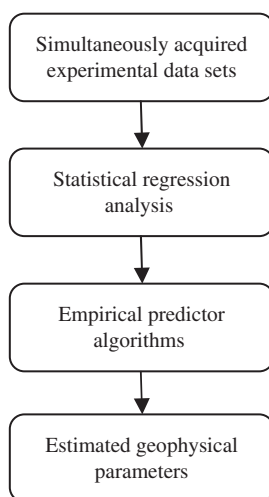


Figure 3. A simplified schematic diagram of the empirical approach for deriving geophysical water-quality parameters from remotely sensed data.

distribution of the parameter from the remotely sensed data. A good example of a commonly used empirical algorithm is the colour-ratio algorithm, which uses a ratio of the reflectance,  $R_i$ , in two spectral channels,  $i$ , to estimate the parameter,  $p$ , where  $\alpha$ ,  $\beta$  and  $\gamma$  are regression coefficients (IOCCG 2000):

$$p = \alpha \left( \frac{R_1}{R_2} \right)^\beta + \gamma. \quad (1)$$

There are many varieties of algorithms that use either single bands, band ratios, band arithmetic or multiple bands as independent variables in linear, multiple linear or non-linear regression analyses (table 1). Once the algorithm has been parameterized, it may be applied to remotely sensed data to estimate the spatial distribution of the parameter in the form of a map. This step usually assumes that the water and atmospheric conditions remain constant throughout the scene. Should the algorithm be applied to scenes acquired at different times or over different areas, the relative errors in the parameter estimates will generally increase as these assumptions break down. The empirical approach when compared to the inverse modelling approach has the key advantages of computational simplicity and ease of implementation. For these reasons, it makes up the majority of studies in inland waters. Normally, empirical algorithms can be expected to perform well only inside their range of derivation and for the area for which they are derived. They are also more limited in their ability to discriminate between non-unique signals from parameters that may be covariant, for example, TSS and Chl-*a*, than inversion algorithms, which solve for a number of parameters simultaneously. Empirical algorithms for estimating  $z_{SD}$ , TSS, Chl-*a*,  $a_{CDOM}$  and other parameters from recent studies in inland and near-coastal transitional waters are reviewed below. The parameters are arranged to reflect the increasing complexity in the methods used. An explanation of the choice of independent variables (bands/band ratios) is often lacking or missing in many empirical studies. This article has attempted to fill this gap by providing brief explanations for the basis of the algorithms in terms of the effect of the IOPs on the remote-sensing reflectance. For other reviews of empirical algorithms, see Kirk (1994), Durand *et al.* (1999), Lindell *et al.* (1999) and (Kutser 2009).

## 4. Review of empirical algorithms

### 4.1 Secchi disk depth or water clarity algorithms

$z_{SD}$  is defined as the depth, in metres, at which a circular white disk disappears when lowered into water as perceived by the human eye and is a measure of the depth of penetration of light in the water or water clarity (Preisendorfer 1986). Secchi disk is inversely proportional to the average amount of inorganic and organic material in the water. Therefore,  $z_{SD}$  is a proxy for gross particulate load or, in humic lakes, the concentration of dissolved organic matter, which is known to have a large effect on water-leaving radiance. Secchi disk derived from remotely sensed data essentially shows changes in gross particulate load, which is useful for obtaining information on biological activity (trophic status) and sediment flux. Thus  $z_{SD}$  is a valuable quantity for aquatic scientists and is also important from a water-management perspective, as water clarity is often the basis with which water users perceive water quality.

There are a large number of studies using Landsat to retrieve  $z_{SD}$ , and most of these use simple linear regressions of single bands or band ratios. In particular, the

use of the red TM3 band is ubiquitous (as single band or ratio) (e.g. Lathrop 1992, Pattiaratchi *et al.* 1994, Cox *et al.* 1998, Härmä *et al.* 2001, Hellweger *et al.* 2004, Kallio *et al.* 2008, Wu *et al.* 2008, Duan *et al.* 2009). In almost all cases, the relationship of  $z_{SD}$  to brightness is non-linear – and therefore  $z_{SD}$  is log transformed in regressions. The correlation with the red band may be causally explained by the direct positive correlation between reflectance in the red and gross particulate load inducing particulate scattering. Therefore, as water clarity ( $z_{SD}$ ) decreases, brightness in the red usually increases. Among these studies, a ratio using the reflectance in the red, TM3, to that in the blue, TM1, is particularly common. Dividing by the blue band, which is dominated by the absorbing effects of phytoplankton, *gelbstoff* and detritus, serves to normalize the brightness in the red band (see figure 1). TM1 is least correlated with  $z_{SD}$  because of the offset to scattering by strong absorption in the blue. A possible exception may be for very clear lakes (e.g. Giardino *et al.* 2001 who used TM1) or for humic lakes – although the very small signal in the blue in these conditions may render it unusable (e.g. Härmä *et al.* 2001). Building on the TM3/TM1 ratio, many studies have subsequently included an additional variable in the form of TM1 (or TM3) in a multiple linear regression to achieve higher correlations (Lavery *et al.* 1993, Kloiber *et al.* 2002, Brezonik *et al.* 2005, Olmanson *et al.* 2008). Thus a multiple regression of the form  $TM3/TM1 + TM1$  may become a common equation for assessing  $z_{SD}$  using Landsat owing to the consistency of the algorithm's performance (Kloiber *et al.* 2002). There are some examples that use green (Lathrop and Lillesand 1986) or NIR bands (SPOT) (Lathrop and Lillesand 1989), although there are few recent examples of this. Less causally explicable multiple regressions with many bands have also been used (Alparslan *et al.* 2007), as well as more advanced approaches such as principal component analysis (Wang and Ma 2001) and artificial neural networks (Zhang *et al.* 2002).

Typical narrow-band ocean-colour sensors such as MODIS and MERIS have also been used successfully to estimate  $z_{SD}$  in lakes and near-coastal transitional waters, with red bands again being used most often. For lakes and coastal waters in Finland, a ratio of the MERIS blue band at 521 nm or 492 nm to the red band at 620 nm or 700 nm proved to give the highest correlations (up to 0.93) (Härmä *et al.* 2001, Kallio *et al.* 2001, Koponen *et al.* 2002, Kratzer *et al.* 2008). The subtraction of an NIR band from each of the bands in some of these studies acts as a rough correction for atmospheric effects. These results again confirm the agreeableness of the blue-to-red ratio for estimating  $z_{SD}$ . It appears that the lack of studies utilizing MODIS to derive water clarity in inland/transitional waters is a result of the relatively coarse resolution of some of the broad visible bands (e.g. bands 3 and 4 are 500 m). However, given the high coefficients of determination observed with the 250 m red band 1 (e.g. Wu *et al.* 2008) and the frequency of acquisition and large quantity of archived data, MODIS has much latent potential to give information on the historical and future trends of water clarity in the world's inland and coastal waters (e.g. Wang *et al.* 2011).

## 4.2 Total suspended solids algorithms

TSS (also called total suspended matter or suspended matter) is the name given to the total mass of suspended particles as measured per volume of water including inorganic (minerals) and organic (detritus and phytoplankton) components. TSS is important for water-quality management, because it is related generally to primary production, sediment transport and, more specifically, water clarity/opacity, which is an indicator of water quality (Dekker *et al.* 2002). It is apparent, however, that far fewer studies

investigate TSS compared with Chl-*a*, and finding empirical algorithms that effectively separate the signals from TSS and Chl-*a* can be challenging.

A difference ratio algorithm of the form  $(R560 - R520)/(R560 + R520)$  was found to be highly correlated with TSS in lakes and rivers not exceeding  $66 \text{ g m}^{-3}$  (Gitelson *et al.* 1993). The algorithm takes advantage of the phytoplankton absorption minimum near 560 nm, which makes the reflectance there sensitive to changes in TSS, while the reflectance at 520 nm is relatively insensitive to changes in TSS (figure 1). Thus the difference ratio acts to normalize the signal at 560 nm for scattering by TSS. More recently, Doxaran *et al.* (2002a) have shown that by exploiting the increased scattering from TSS in the NIR in the highly turbid ( $<985 \text{ g m}^{-3}$ ) waters of the Gironde Estuary, France, a ratio of the reflectance near 850 nm to that at about 550 nm also exhibits a strong non-linear correlation. This result shows the high potential of NIR bands for TSS estimations, as will be shown later on in the discussion. TSS has also been estimated using the reflectance peak near 700 nm, much like Chl-*a* from sensors such as MERIS in relatively low concentrations  $<32 \text{ g m}^{-3}$  ( $r^2 > 0.81$ ) (Härmä *et al.* 2001, Kallio *et al.* 2001, Koponen *et al.* 2007). However, this may give rise to significant ambiguity between TSS and Chl-*a* estimations from satellite data, especially when they are, as they often are, significantly covariant.

The retrieval of TSS from broad-band sensors (including the broad MODIS 250 m bands 1 and 2) has in general been more successful than that of Chl-*a* in terms of significance of correlations (Lathrop and Lillesand 1989, Dekker *et al.* 2002, Doxaran *et al.* 2002b, 2006, 2009, Onderka and Pekarova 2008, Petus *et al.* 2010). In many instances, simple linear regressions of single bands and band ratios give sufficiently good correlations ( $r^2 > 0.82$ ) (Östlund *et al.* 2001, Doxaran *et al.* 2002b, 2006, 2009, Miller and Mckee 2004, Sváb *et al.* 2005, Tyler *et al.* 2006, Onderka and Pekarova 2008). However, there is no immediately apparent agreement between studies reviewed here on which bands/ratios are best for TSS estimation. Lathrop and Lillesand (1989) found that, at high concentrations ( $>10 \text{ g m}^{-3}$ ), the red (XS2) and NIR (XS3) bands of SPOT 1 HRV are more sensitive to changes in TSS than at low concentrations, where a ratio of red to green (XS2/XS1) proved more sensitive. Thus they used a multispectral approach by combining the red-to-green ratio and the NIR band to account for the entire data range ( $4.6\text{--}28.9 \text{ g m}^{-3}$ ,  $r^2 = 0.93$ ). The usefulness of single red and NIR bands has been confirmed over similar concentrations by other studies, including those using the MODIS 250 m bands 1 (620–670 nm) and 2 (841–876 nm) (e.g. Miller and Mckee 2004, Tyler *et al.* 2006, Onderka and Pekarova 2008, Petus *et al.* 2010). More recent work by Nechad *et al.* (2010) used bio-optical theory to prove the usefulness of single red or NIR band algorithms for estimating TSS in turbid waters from MERIS, MODIS and SeaWiFS. The proof of concept of using the single visible band (600–700 nm) of the geostationary SEVIRI on board METEOSAT Second Generation to estimate TSS has already been confirmed (Neukermans *et al.* 2009, Salama and Shen 2010). The reason for the correlation is explained by the contribution of particulate matter, particularly the inorganic component, to scattering in the red and NIR.

However, Dekker *et al.* (2002) pointed out that, for Landsat, the sensitivity is lower, the radiometric calibration is less reliable and the influence of the adjacency effect is more severe in the NIR, citing potential drawbacks of using a single Landsat TM NIR band (see also Ruddick *et al.* 2006). Nonetheless, if these problems can be solved through new more advanced sensors and better atmospheric correction procedures, then single-band NIR algorithms would be very effective for estimating TSS. Using a



bio-optical modelling approach, Dekker *et al.* (2002) found that an exponential relationship between the average of the green and red bands of Landsat TM (TM2, TM3) and SPOT 3 HRV (XS1, XS2) was highly correlated ( $r^2 = 0.99$ ) with TSS in the range  $5\text{--}50\text{ g m}^{-3}$  for turbid lakes in the Netherlands. This algorithm takes advantage of the reflectance maximum near 560 nm in the green (explanation given above) and the increasing reflectance in the red, and avoids the current problems with using single NIR bands. In estuarine waters with very high concentrations of TSS ( $>2000\text{ g m}^{-3}$ ) Doxaran *et al.* (2002b, 2003, 2006) showed, through using a bio-optical model, the robustness of the NIR-to-green and NIR-to-red band ratios for estimating TSS from SPOT over long timescales. The use of band ratios, as opposed to the single NIR band, normalizes (reduces) the effects of variable sediment refractive indices and particle grain sizes. Thus there may be advantages using ratios over single bands in instances where algorithms are expected to perform well over large variations in TSS caused by tidal or seasonal variations, such as occurs in estuaries (e.g. Doxaran *et al.* 2009). From the above examples, it would appear that the choice of algorithm for TSS when using broad-band sensors largely depends on the concentration ranges and area of investigation.

An alternative approach to estimating TSS is to use multiple linear regressions (e.g. Wang and Ma 2001, Wang *et al.* 2006, Alparslan *et al.* 2007). Although these procedures may give high coefficients of determination, they lack usually causal explanations and cross-temporal applicability of those reviewed already. More advanced-type algorithms such as artificial neural networks (discussed further below) using multiple bands have also been used to improve retrieval accuracies (Zhang *et al.* 2002, Sudheer *et al.* 2006).

### 4.3 Phytoplankton pigment algorithms (Chl-*a*)

Phytoplankton pigment concentration (Chl-*a*) is the most commonly derived parameter in water-quality remote sensing mainly because of its use in determining the trophic status of waters. It acts as a proxy for phytoplankton concentration and is therefore an important component in the derivation of secondary products such as primary production.

**4.3.1 R700/R670-based colour-ratio algorithms.** The ratio of reflectance at about 700 nm to that near 670 nm has been widely used for estimating Chl-*a* concentration in high-biomass waters (see table 1). In particular, studies by Mittenzwey *et al.* (1992), Gitelson *et al.* (1993), Gons (1999) and more recently Moses *et al.* (2009a) show that the correlation between Chl-*a* concentration and the ratio is very significant ( $r^2 > 0.8$ ) for a variety of waters, including rivers, lakes, estuaries and in the laboratory, and over a wide range of concentrations from  $\sim 0.1$  to  $350\text{ mg m}^{-3}$ . There are a large number of studies that use high spectral resolution airborne and *in situ* radiometric data (Dierberg and Carriker 1994, Jiao *et al.* 2006, Duan *et al.* 2007, Hunter *et al.* 2008b, Hunter *et al.* 2009) to estimate Chl-*a* using the ratio, and the  $r^2$  values for these studies range from 0.75 (Duan *et al.* 2007) to 0.99 (Menken *et al.* 2006). The positioning of the MERIS bands at 665 and 709 nm makes MERIS ideally suited for predicting Chl-*a* using this ratio, and many studies have recently been carried out (Flink *et al.* 2001, Ammenberg *et al.* 2002, Härmä *et al.* 2001, Kallio *et al.* 2001, Gons *et al.* 2002, Koponen *et al.* 2002, Kallio *et al.* 2003, Strömbeck *et al.* 2004, Gons *et al.* 2005, Koponen *et al.* 2007, Moses *et al.* 2009b) where the coefficients of determination range from 0.84 (Flink *et al.* 2001) to 0.98 (Kallio *et al.* 2003). These

studies generally use relatively simple linear, multiple linear and non-linear power-law regression or polynomial analyses.

The causal explanation for the strength of the correlation of Chl-*a* with the 700/670 nm ratio is based on the interaction between backscattering from particulate matter (phytoplankton) and the strong absorption of water, which both increase towards the infrared (figure 1). The offset to scattering due to absorption by water near 700 nm causes a sharp peak in highly scattering (turbid or productive) waters. The height and position of the peak is known to be well correlated with Chl-*a*, with the peak shifting towards greater wavelengths ( $\sim 715$  nm) as Chl-*a* increases (Gitelson 1992). In contrast, the reflectance near 670 nm is uncorrelated, with Chl-*a* being almost constant owing to the Chl-*a* absorption maximum, which offsets backscattering. Thus the 700/670 nm ratio can be effectively exploited to determine Chl-*a*, as it normalizes the signal from particulate phytoplankton backscattering.

There are a number of studies that employ slight variations on the ratio. By subtracting an NIR band (781 or 754 nm) from the 709 nm and 665 nm bands as a rough atmospheric correction, some of these studies improve the strength of the correlation (e.g. Härmä *et al.* 2001, Koponen *et al.* 2002). This is based on the assumption that the reflectance in the NIR over water is mainly due to atmospheric effects, which may be a false assumption over turbid water. Some studies make use of the height of the maximum reflectance peak near 700 nm ( $R_{\max}/R_{670}$ ) rather than a fixed bandwidth, and have achieved high correlations ( $r^2 = 0.95$ ) for a wide range of concentrations in Lake Kinneret, Israel (Yacobi *et al.* 1995). Others have used the peak near 560 nm rather than at 709 nm in low biomass conditions (Floricioiu *et al.* 2004, Candiani *et al.* 2005). Gons (1999) and Gons *et al.* (2002, 2005) restated the ratio in terms of the commonly used reflectance approximation (Morel and Prieur 1977) and used *in situ* reflectance and Chl-*a* data to calibrate the value of the Chl-*a* specific absorption coefficient,  $a_{\phi}^*$ , at 672 nm. This algorithm makes a number of assumptions, including that the backscattering is spectrally invariant and can be sufficiently estimated from the NIR wavelengths. Nevertheless, the algorithm provides good results in well-mixed, high-biomass waters. A more pronounced variation on the ratio is the so-called 'three-band model' proposed by Gitelson *et al.* (2003):

$$\text{Chl-}a = R(\lambda_3) \left[ \frac{1}{R(\lambda_1)} - \frac{1}{R(\lambda_2)} \right] \quad (\text{mg m}^{-3}), \quad (2)$$

where  $R(\lambda_1)$  is reflectance in wavelength maximally sensitive to Chl-*a* absorption (670 nm),  $R(\lambda_2)$  is reflectance in wavelength minimally sensitive to absorption by Chl-*a* (710 nm) and  $R(\lambda_3)$  is reflectance in wavelength minimally effected by absorption that accounts for scattering (750 nm).

The three-band algorithm has been used to give very good estimates of Chl-*a* in turbid and very high biomass hypertrophic waters (Dall'Olmo and Gitelson 2005, Zimba and Gitelson 2006, Gitelson *et al.* 2008, 2009, Moses *et al.* 2009b). A new four-band algorithm, including an additional band near 700 nm, was found to be an improvement over the three-band model in highly turbid lake water through better accounting for absorption by water and non-negligible scattering by suspended matter in the NIR (Le *et al.* 2009).

**4.3.2 The fluorescence/reflectance line height algorithm.** The fluorescence maximum near 685 nm has often been used to estimate Chl-*a* (Gower 1980, Gitelson *et al.* 1994). The fluorescence line height (FLH) algorithm measures the height of the fluorescence



peak at 685 nm from a linear baseline drawn between two points on either side of the peak (Dierberg and Carriker 1994, Giardino *et al.* 2005). The coefficients of determination for linear regression FLH algorithms range from 0.73 (Gitelson *et al.* 1994) to 0.86 (Dierberg and Carriker 1994) for Chl-*a* concentrations not exceeding 79 mg m<sup>-3</sup> in this instance. It is important to consider that the FLH algorithm is only suitable for use in relatively low-biomass waters (Chl-*a* concentrations generally not exceeding ~30 mg m<sup>-3</sup>) as the backscattering peak near 700 nm overwhelms the fluorescence peak in high-biomass water. Therefore, it is very difficult, or impossible, to differentiate between the signal from particulate backscattering and solar-induced fluorescence in high-biomass waters. A very similar algorithm that is better suited to high-biomass waters is the reflectance line height (RLH) or scattered line height algorithm (Dierberg and Carriker 1994, Yacobi *et al.* 1995, Schalles *et al.* 1998). The RLH algorithm is a modified FLH algorithm and is written (after Gower *et al.* 1999) as follows:

$$RLH = L_2 - L_1 - \left[ \frac{(L_3 - L_1) * (\lambda_2 - \lambda_1)}{(\lambda_3 - \lambda_1)} \right], \quad (3)$$

where  $L_i$  is the radiance in band  $i$  and  $\lambda_i$  is the centre wavelength of band  $i$ . Band  $i = 2$  is centred on the reflectance peak close to 700 nm, whereas bands  $i = 1, 3$  on either side determine the baseline (670 and 850 nm).

In waters with high Chl-*a* concentrations,  $r^2$  values range from 0.85 (Dierberg and Carriker 1994) to 0.96 (Yacobi *et al.* 1995). The maximum chlorophyll index or MCI for MERIS is derived using the RLH algorithm and the 709 nm band (Gower *et al.* 2005). A minor variation on the RLH algorithm is the SUM algorithm, which uses the sum of the area under the reflectance peak using a baseline between 670 nm and 730 or 850 nm (see Gitelson *et al.* 1994, Schalles *et al.* 1998). The SUM algorithm gave  $r^2$  values >0.84 for turbid productive waters (Schalles *et al.* 1998).

**4.3.3 Broad-band algorithms.** Sensors such as Landsat, SPOT and IKONOS are often used to detect Chl-*a* (e.g. Lathrop and Lillesand 1986, Dor and Ben-Yosef 1996, Thiemann and Kaufmann 2000, Giardino *et al.* 2001, Nas *et al.* 2009, Ormeci *et al.* 2009; table 1). However, the lack of narrow bands and low SNRs prevents the use of the more specialized algorithms already described. Therefore, simple linear regressions of single bands or band ratios (which may be log-transformed) are often used and occasionally give coefficients of determination as high as 0.98 (Lathrop and Lillesand 1986), but more often give less-significant correlations. There is little consistency between the studies concerning which band ratios or bands are useful to detect Chl-*a*, with bands/ratios often selected in a seemingly haphazard manner. With a few exceptions (e.g. Gitelson *et al.* 1996, Brivio *et al.* 2001), there is often also no attempt to explain the observed correlation between bands/band ratios and Chl-*a* causally. A ratio using the reflectance in the red band (TM band 3) and that in the blue (TM band 1) may be useful in this regard, as Chl-*a* is directly proportional to the magnitude of the reflectance in band 3 due to the influence of the reflectance peak in the red, and inversely proportional to that in the blue, as a consequence of the chlorophyll absorption maximum in this wavelength (Gitelson *et al.* 1993, Gitelson *et al.* 1996, Han and Jordan 2005). Similarly, a ratio using the red band and NIR band (TM band 4) gives some consistent results as the reflectance in the NIR may sometimes be assumed to be close to zero and less variable due to strong absorption by water (Yacobi *et al.* 1995, Duan *et al.* 2007). At lower Chl-*a* concentrations (<20 mg m<sup>-3</sup>), it appears that the ratio of the green band (TM band 2) to the blue may be better

suited, as the correlation with Chl-*a* shifts to slightly lower wavelengths in the red (Gitelson *et al.* 1993, Östlund *et al.* 2001). Another approach that gives good correlations in these conditions is to subtract the reflectance in the red from that in the blue to remove effects of scattering by inorganic suspended solids and normalize using the green reflectance peak band (Mayo *et al.* 1995, Brivio *et al.* 2001). Normalizing the reflectance in a band by integrating the reflectance in all visible bands (or the sum of the bands) may also lead to improved correlations (Gitelson *et al.* 1993). The alternative approach to these ratios is to use multiple linear regressions combining a number of bands or ratios. However, a causal explanation is deficient and the cross applicability of these algorithms is normally very limited. Multiple variable linear regressions may use as many as four bands to give very high correlations (e.g. Giardino *et al.* 2001, Nas *et al.* 2009, Ormeci *et al.* 2009, table 1). More advanced and creative methods are then needed to establish causality and improve the cross-applicability of the procedures.

**4.3.4 Advanced algorithms.** Advanced algorithms are often employed in order to improve the strength of correlations and the cross-spatial/temporal applicability of algorithms. These include artificial neural network algorithms (Zhang *et al.* 2002, Sudheer *et al.* 2006, Wu *et al.* 2009), genetic algorithms (Chen *et al.* 2008, Wu *et al.* 2009), multivariate regression analysis using classification procedures such as the linear mixture modelling approach (Tyler *et al.* 2006) and spectral decomposition algorithm (Oyama *et al.* 2007, 2009). Of these, the neural network approach has become particularly popular. Neural networks are essentially a multiple non-linear regression procedure simulating the neural functions of the cerebral cortex of the brain through layers of interacting neurons. The ‘input layer’ uses the reflectance/radiance in various bands and the ‘output layer’ gives the concentrations of various parameters or IOPs. The hidden layers consist of neurons that are assigned various weights and bias through training the algorithm with a real or simulated data set of concentrations and reflectances. Neural networks are able to simulate the complex non-linear influence of various parameters on the water-leaving reflectance and solve for a variety of parameters simultaneously. For this reason, they are also used for analytical algorithms. However, the need for a training data set and problems such as ‘over-training’ and multiple possible solutions means that these algorithms require substantial effort to implement. Genetic algorithms simulate the process of natural evolution of populations through producing successively ‘fitter’ generations through processes of selection, crossover and mutation. These algorithms have a powerful ability to search the full decision space for the optimal solution to linear and non-linear problems but are often combined with procedures such as neural networks that improve the precision of the solution (e.g. Wu *et al.* 2009). The use of these and other complex algorithms generally leads to improved significance of correlations, although, in some instances, the level of complexity of these algorithms does not appear to offer significant enough accuracy improvements to justify their mathematical difficulty.

#### 4.4 CDOM algorithms

CDOM (also called *gelbstoff*, *gilvin* or yellow substances), composed of humic and fulvic acids, is a significant contributor to water colour, because humic substances absorb strongly in the blue region of the spectrum turning the water brown. Absorption by CDOM ( $a_{\text{CDOM}}$ ), usually referenced at 440 nm, takes on the form of an

exponential curve decreasing towards longer wavelengths so that its effects are usually negligible at wavelengths  $> \sim 550$  nm. The slope of the curve is mostly predictable, as it varies within a relatively small range ( $0.10$ – $0.20$  nm $^{-1}$ ) for most inland and coastal waters (Dekker 1993). Absorption by detrital material (organic non-living material of mostly algal origin), mineral and non-pigmented aquatic particles display similar decreasing exponential functions (Babin and Stramski 2002, Babin *et al.* 2003) and can overwhelm the contribution to absorption by CDOM in natural waters if concentrations are high enough. In turn, in situations where  $a_{\text{CDOM}}$  is very large, that is, humic lakes, the retrieval of other biogeophysical parameters will also be affected.

The retrieval of  $a_{\text{CDOM}}$  using remote sensing has been the subject of more recent studies, as interest in optical measurements of the IOPs of natural waters is increasing (Lee *et al.* 2002). Absorption by CDOM is one of the primary additive absorption IOPs, along with phytoplankton and water, and is therefore of great interest from a bio-optical perspective. Because the signal from  $a_{\text{CDOM}}$  is only significant in the blue region of the spectrum ( $< 550$  nm), it serves to reason that the retrieval from remote sensing will utilize suitable bands from this region of the spectrum. An unfortunate consequence of this is that atmospheric scattering is greatest in the blue – and the water-leaving signal so diminished from strong absorption by phytoplankton and CDOM – that the signal from the water may be indistinguishable and the data unusable. This is especially a problem when considering the low radiometric sensitivities of sensors such as Landsat or SPOT. Despite these limitations, there are numerous examples of  $a_{\text{CDOM}}$  retrieval using both sensitive ocean colour and radiometric sensors and less-sensitive sensors, albeit in some cases with the acceptance of lower correlations.

Algorithms using ratios of reflectance in the blue ( $\sim 400$ – $500$  nm) to that in the green or red ( $\sim 500$ – $700$  nm) have been found to be well correlated with  $a_{\text{CDOM}}$  (Kutser *et al.* 1998). In waters with low suspended sediment, Bowers *et al.* (2000) showed theoretically, while making some assumptions about particulate absorption, that there is a linear relationship between  $a_{\text{CDOM}}$  and the ratio of reflectance in the red to the blue. This was confirmed by a small data set ( $n = 8$ ) using the 670/412 nm ratio ( $r^2 = 0.99$ ) and other studies (e.g. Bowers *et al.* 2004). In turbid waters, however, this relationship breaks down due to the influence of particulate scattering (Fang *et al.* 2009) and is therefore better described by a non-linear power-law relationship. Doxaran *et al.* (2005) used a 400/600 nm ratio, whereas D'Sa and Miller (2003) used the SeaWiFS band configurations 412/510, 443/510 and 510/555 nm, all of which gave good results, although this may reflect the existence of strong covariance between Chl-*a* and CDOM. Comparable red/blue ratios produced with the MERIS waveband configuration also give similarly strong correlations up to 0.96 (Kallio *et al.* 2001, Ammenberg *et al.* 2002, Koponen *et al.* 2007). Ratios used include MERIS bands 665/490 nm and 665/550 nm. The use of the 665 nm band takes advantage of the Chl-*a* absorption maximum and acts to normalize for the effects of Chl-*a* absorption and backscattering by particulate matter. The correlation with the 571–607/607 nm ratio used by Kallio *et al.* (2001) may, however, be as a result of covariance between TSS and  $a_{\text{CDOM}}$ , and is therefore less likely to be applicable elsewhere as the signal from CDOM at 570 nm is small.

To discriminate the signal from phytoplankton pigment absorption and that from CDOM in the blue more effectively, Gitelson *et al.* (1993) proposed a 'decoding index' difference ratio algorithm to retrieve  $a_{\text{CDOM}}$  from high-resolution radiometric data, where  $a$  and  $b$  are regression coefficients:

$$a_{\text{CDOM}} = a \left[ \left( R_{480} - \frac{R_{700}}{R_{675}} - R_{520} \right) / \left( R_{480} + \frac{R_{700}}{R_{675}} + R_{520} \right) \right]^b \quad (\text{m}^{-1}). \quad (4)$$

The 480 nm band is strongly influenced by  $a_{\text{CDOM}}$ , whereas the band at 520 nm is a reference and the  $R_{700}/R_{675}$  ratio a correction factor for Chl-*a* absorption. The non-linear power-law algorithm gave an  $r^2$  value of  $>0.9$  for  $a_{\text{CDOM}}$  at 380 nm in the range  $0.1\text{--}12 \text{ m}^{-1}$  in  $>20$  inland water bodies.

Broad-band terrestrial sensors can also be used successfully to estimate  $a_{\text{CDOM}}$  in lakes and transitional waters, although usually not with the same level of accuracy as with sensitive and high spectral definition instruments. As a result of atmospheric interference, very small signal and influences of Chl-*a* absorption, the use of a broad band in the blue (TM1) is limited or infeasible (Kutser *et al.* 2005b, Kallio *et al.* 2008). Thus CDOM is better estimated taking advantage of its inverse correlation with the band in the green: as CDOM increases, the reflectance in the green gradually decreases (Kutser *et al.* 2005, Kallio *et al.* 2008). Thus, a ratio of the green to red (where influence by CDOM is assumed to be zero) has been shown to be the most useful algorithm (Kutser *et al.* 2005a, 2009). This may mean that detecting CDOM by terrestrial sensors at low concentrations ( $<\sim 5 \text{ m}^{-1}$  at 440 nm) that do not have a significant impact on reflectance  $>500 \text{ nm}$  is impossible. However, even when using a green/red algorithm with large  $a_{\text{CDOM}}$ , the low radiometric resolution of some sensors makes CDOM estimations infeasible (Kutser *et al.* 2005b). Results show that the minimum acceptable radiometric resolution for mapping CDOM from broad-band sensors is about 16 bit – making the TM and IKONOS instruments generally ill-suited (Kutser *et al.* 2005b). Nevertheless, successful  $a_{\text{CDOM}}$  estimations have been given from TM (8 bit) (Kallio *et al.* 2008) and ALI (16 bit) (Kutser *et al.* 2005a, b, Kutser *et al.* 2009) in high CDOM boreal lakes and coastal waters in Finland. These used a non-linear power-law regression of the green/red band ratio to give  $r^2$  values  $<0.84$ . Using multiple linear regressions, Brezonik *et al.* (2005) derived  $a_{\text{CDOM}}$  from TM with an  $r^2$  value of 0.77 for high CDOM lakes in Minnesota.

#### 4.5 Other parameters algorithms

A wide variety of additional parameters has also been retrieved in lakes and near-coastal transitional waters – some of which are included in the studies presented in table 1. Some of these parameters, such as nutrient concentrations of phosphorus or nitrogen, are derived via proxy using their relationship to parameters that are optically active. These, however, are not reviewed here: only parameters that are directly attainable due to their effects on the water-leaving radiance are included. These are, most notably, turbidity, suspended particulate inorganic matter (SPIM) and PC, the accessory pigment present in blue-green algae.

**4.5.1 Turbidity.** Turbidity is closely related to TSS and  $z_{\text{SD}}$ , measuring either the attenuation of a beam of light passing through water or, more commonly, the scattering by particulate matter usually as detected within  $90^\circ$  of the light beam [units = nephelometric turbidity units (NTU)]. The relationship between TSS and turbidity is variable in space and time owing to variations in particle size distribution, shape, composition and reflective index (Chen *et al.* 2007, Kallio *et al.* 2008). There are numerous recent examples of remote estimation of turbidity from MODIS 250 m (Chen *et al.*

2007, Alcântara *et al.* 2009, Petus *et al.* 2010), Landsat (Lathrop and Lillesand 1986, Vincent *et al.* 2004, Brezonik *et al.* 2005, Wang *et al.* 2006, Kallio *et al.* 2008), other broad-band sensors (Lathrop and Lillesand 1989, Hellweger *et al.* 2007, Chen *et al.* 2009) and others (Dierberg and Carriker 1994, Fraser 1998, Koponen *et al.* 2002). As for TSS, algorithms using single bands or ratios in the red are generally most successful owing to the influence of particulate scattering in these bands, provided the impact from phytoplankton pigments is not overwhelming (Kallio *et al.* 2008). Some studies use more advanced spectral unmixing (Alcântara *et al.* 2009) and NN algorithms (Zhang *et al.* 2002).

**4.5.2 The inorganic component of suspended matter.** SPIM may be of interest when large amounts of sediment are entrained in the water column such as in tidal estuaries (e.g. Doxaran *et al.* 2005). Algorithms for SPIM detection are very similar to those for TSS (especially when inorganic matter makes up the majority of TSS) and use red and NIR bands (e.g. Ammenberg *et al.* 2002, Doxaran *et al.* 2005). This is because inorganic particles contribute greatly to scattering in the NIR (Doxaran *et al.* 2007).

**4.5.3 Phycocyanin.** PC is an accessory pigment present in certain species of cyanobacteria. The detection of cyanobacteria via PC is of great interest owing to the potentially great negative impacts these bloom-forming and sometimes toxic species have in coastal and inland waters, which may result in substantial economic losses. Therefore, an effective means of detecting and monitoring these blooms would be of considerable public value. There have already been numerous attempts to develop an empirical algorithm specifically for detecting PC; there have been other studies aimed at detecting cyanobacterial blooms via Chl-*a* (e.g. Kutser 2004, Reinart and Kutser 2006). Absorption by PC  $\sim 620$  nm in turbid cyanobacteria-dominated water results in a distinct minimum in the water-leaving reflectance, which can potentially be used to detect cyanobacteria-specific blooms (Dekker 1993, Jupp *et al.* 1994). Therefore, there should be an inverse relationship between the reflectance near 620 nm and PC concentration – although the influence of water, *tripton*, other phytoplankton pigments and to a lesser extent CDOM can be a significant source of error owing to the relative magnitudes of these absorption coefficients. Furthermore, empirical algorithms exploiting this unique feature  $\sim 620$  nm require sensors with narrow bands ( $<12$  nm wide – conveniently MERIS has 10 nm wide bands centred at 620 nm) and high SNRs ( $>1000:1$ ) (Dekker 1993, Metsamaa *et al.* 2006). Single bands ( $\sim 620$  nm) and the band ratio 620/650 nm have been used (Dekker *et al.* 1992, Gitelson *et al.* 1995, Schalles and Yacobi 2000, Ruiz-Verdú *et al.* 2008, Vallely 2008, Mishra *et al.* 2009) but are quite susceptible to errors arising from spectral variations of extraneous parameters. To avoid this problem, Dekker (1993) subtracted the reflectance at 624 nm from the average of that at 600 and 648 nm as a baseline subtraction and found a strong linear relationship with PC for eutrophic turbid lakes ( $r^2 > 0.99$ ). However, further study shows that the algorithm has a tendency to underestimate and may only be sensitive at concentrations  $>\sim 10$  mg m $^{-3}$  and  $<\sim 200$  mg m $^{-3}$ , as the relationship with the 620 nm reflectance trough is non-linear outside these ranges (Ruiz-Verdú *et al.* 2008).

Other studies have used the 709/620 nm ratio with success (Peña-Martínez *et al.* 2004, Ruiz-Verdú *et al.* 2005, Hunter *et al.* 2008b, Hunter *et al.* 2009) or a 700/600 nm ratio (Mishra *et al.* 2009). To account for the complex interactions of absorption by water and Chl-*a* (but not CDOM/detritus), Simis *et al.* (2005) and Simis *et al.* (2007)



used a semi-analytical algorithm based on the reflectance approximation (Morel and Prieur 1977) and the 709/620 nm ratio to retrieve the PC-specific absorption coefficient at 620 nm,  $a_{pc}(620)$ :

$$a_{pc}(620) = \left\{ \left[ \frac{R(709)}{R(620)} \right] [a_w(709) + b_b] \right\} - b_b - a_w(620) \delta^{-1} - [\epsilon a_{chl}(665)] \quad (\text{m}^{-1}), \quad (5)$$

where  $a_w$  is the absorption by water,  $b_b$  is the spectrally invariant backscattering coefficient,  $\delta$  is a correction factor that accounts for the assumption that absorption by CDOM, Chl-*a* and detritus is negligible in the red,  $a_{chl}$  is the absorption due to chlorophyll and  $\epsilon$  is a conversion factor to estimate  $a_{chl}$  at 620 nm from that at 665 nm.

The algorithm uses  $b_b$  calculated from reflectance at 778 nm according to Gons (1999), and  $a_{chl}(665)$  calculated from a very similar semi-analytical algorithm using the 709/665 nm ratio (Simis *et al.* 2005). Final PC concentrations are obtained by dividing  $a_{pc}(620)$  by the PC-specific absorption coefficient  $a_{pc}^*(620)$ . This algorithm has been validated by a number of other studies ( $r^2 < 0.99$ ) (Simis *et al.* 2007, Randolph *et al.* 2008, Ruiz-Verdú *et al.* 2008, Vallely 2008) and similar semi-analytical algorithms (Yang and Pan 2006). The proven cross-applicability of such an approach makes it potentially very useful for PC estimations in cyanobacteria-dominant eutrophic waters.

As previously discussed, sensors with high spatial resolution, low SNR and wide bandwidths do not meet the minimum requirements for PC detection. Nonetheless, using Landsat TM and ETM+, Vincent *et al.* (2004) found that multiple linear regressions of band ratios were capable of establishing a correlation with PC ( $r^2 < 0.78$ ). However, the causality and cross-applicability of such an approach is very limited especially given that the Landsat bands do not overlap the PC absorption maximum at 620 nm. The result is probably explained by a correlation between PC and turbidity or some other parameter (Kutser *et al.* 2006). In general, PC retrieval is best when cyanobacteria form the dominant component of the phytoplankton assemblage due to the confounding effects of other accessory pigments from other phytoplankton groups (Ruiz-Verdú *et al.* 2008). PC estimation from remotely sensed data such as MERIS and new hyperspectral space-borne sensors (e.g. HICO, Enmap hyperspectral Imager) will likely be used in future to assess cyanobacterial blooms and provide early-warning systems (Ahn *et al.* 2007).

## 5. Discussion and conclusion

Progress towards a bio-optical model-based approach of remote sensing of water has been a necessary advancement of the science of what is now rather inappropriately called ocean-colour radiometry. This shift has been caused by the need for a technique capable of solving the complex signal from mainly coastal waters. Model-based solutions have the ability to retrieve simultaneously various optical and biogeophysical parameters from the remote-sensing reflectance. Their theoretical basis on sound solutions to the radiative transfer equation makes them potentially a more expedient approach. So, why write a review focusing on the empirical approach when the science is clearly heading towards model-based approaches? First, there is a growing amount of evidence from recent studies that the empirical approach is more often used than the model-based approach and is fully capable of producing reliable estimates of a

variety of biogeophysical parameters in inland and transitional waters. Furthermore, the ease of implementation and computational simplicity often makes it the most convenient choice for many water remote-sensing studies. These advantages are also favourable for near real-time observation systems, especially where the deployment of autonomous observation moorings/platforms presents a readily available source of data that can easily be used to parameterize and validate empirical algorithms.

Second, empirically based algorithms have long been used to derive standard products for phytoplankton pigments (Chl-*a*) in open-ocean waters, for example, the SeaWiFS OC4.v4 (O'Reilly *et al.* 1998) or MODIS OC3M (Campbell and Feng 2005) standard algorithms. These algorithms have been parameterized and validated using the great amount of *in situ* data collected with considerable effort over several decades [stored in databases, e.g. the SeaWiFS Bio-Optical Archive and Storage System (SeaBASS) and the NASA bio-optical Marine Algorithm Data set (NOMAD), the new MERIS Matchup *In-situ* Database (MERMAID)]. These empirical algorithms for the open ocean are limited in coastal waters, hence the need for a model-based approach. However, what about empirical algorithms derived specifically for these coastal, inland and transitional waters? Following the review of the large number of studies, the application of empirically based algorithms to produce common biogeophysical products for inland and transitional waters appears achievable. For example, the 709/664 ratio is remarkably effective for detecting high-biomass waters typical of eutrophic/hypertrophic systems (see discussion below). Until now, only very limited attempts have been made to produce common biogeophysical algorithms in complex coastal waters, lakes or estuaries (e.g. Nechad *et al.* 2003, Gower *et al.* 2005). Parameterization and validation of such algorithms could be achieved using the great amount of *in situ* data that have been collected in recent years and will in future be collected from autonomous platforms. This could lead to the development of regionally optical water type-specific and perhaps even, for certain applications, global empirically based algorithms, such as the MCI (Gower *et al.* 2008), although the latter remains speculative at present.

Third, the bio-optical basis and potential to operate 'independently' of *in situ* data, often cited as the main advantages of model-based algorithms, is also true for some empirical algorithms. For example, studies such as Bowers *et al.* (2000), Dekker *et al.* (2001, 2002) and Doxaran *et al.* (2006) used bio-optical modelling to derive robust *empirical* algorithms for estimating  $a_{CDOM}$ , TSS in Dutch lakes and TSS in the Gironde estuary, France, respectively. These algorithms are in theory able to operate independently from *in situ* data and have the strong theoretical basis of model-based algorithms. There is a need for more studies investigating bio-optical modelling for generating empirical algorithms. It should also be noted that, in reality, a great deal of *in situ* data on IOPs are required for parameterization and validation of model-based algorithms if they are to operate reliably on a regional or even local basis. The evidence given above in support of the empirical approach shows that attempts to review and further develop these techniques is justified. Table 3 presents a synthesis of the findings in section 4.

Table 3 can be used as a guide for selecting appropriate empirical algorithms for use with either broad- or narrow-band sensors in various inland and transitional waters. It presents suitable detection band(s), band ratios and band arithmetic-independent variables for different concentration ranges of the parameters. The explanation of the bio-optical basis establishes a link between the independent variables and the parameter of interest, which is important for the cross-applicability and reproducibility of prospective algorithms. From the information in table 3, empirical algorithms may be created that have a wide range of potential applications for satellite as well as *in situ*

Table 3. Suggested band(s), band ratios and/or band arithmetic for the detection of water-quality parameters in inland and transitional waters using broad-band or narrow-band sensors based on review of current literature.

Parameter	Sensor spectral resolution		Bio-optical basis
	Broad bands	Narrow bands (nm)	
$z_{SD}$	Red band or red/blue ratio, e.g. TM3/TM1 + TM1	Red band or blue/red ratio, e.g. 512/620	Reflectance in red $\propto b_{bp}$ . The blue band dominated by $a_\phi$ and $a_{CDOM}$ serves to normalize
TSS	$<10 \text{ g m}^{-3}$ : Red/green ratio or (green + red)/2	$<30 \text{ g m}^{-3}$ : (560–520)/(560 + 520) or single red band, e.g. 700	The $a_\phi$ minimum at 560 nm is sensitive to TSS, whereas the 520 nm band serves to normalize
	$>10 \text{ g m}^{-3}$ : Red or NIR band or (green + red)/2	$>30 \text{ g m}^{-3}$ : NIR ratio, e.g. 850/550	Reflectance in red and NIR $\propto b_{bp}$ and $b_{bm}$ . Band ratios normalize for variations in particle refractive indices and grain sizes
	$>30 \text{ g m}^{-3}$ : NIR/red or NIR/green ratio		
Chl- <i>a</i>	$<20 \text{ mg m}^{-3}$ : Green/blue ratio or (blue – red)/green	$<30 \text{ mg m}^{-3}$ : 560 or FLH algorithm	Chl- <i>a</i> $\propto$ reflectance in red due to $b_{bp}$ , and inversely related to reflectance in blue due to $a_\phi$
	$>20 \text{ mg m}^{-3}$ : Red/blue or red/NIR ratio	$>30 \text{ mg m}^{-3}$ : 700/670 ratio or three-band model 750(1/670–1/710) or RLH or SUM algorithms	Reflectance at 700 nm sensitive to $b_{b\phi}$ normalized by the $a_\phi$ maximum near 665 nm
$a_{CDOM}$	Green/red ratio	Red/blue ratio, e.g. 670/412, or ‘decoding index’ [490–(700/675)–520]/[490 + (700/675)+ 520]	Relatively insensitive sensors: Reflectance in green inversely related to $a_{CDOM}$ normalized by reflectance in red  Sensitive sensors: Reflectance in blue inversely related to $a_{CDOM}$ normalized by the reflectance in the red
Turbidity	Red band	Red or NIR band	Reflectance in red and NIR $\propto b_{bp}$ and $b_{bm}$
SPIM	Red or NIR band	Red or NIR band	Reflectance in red and NIR $\propto b_{bp}$ and $b_{bm}$
PC	–	(620/650) or (709/620) ratio or (600 + 648)/2–624	Reflectance at 620 nm inversely related to PC due to absorption maximum

Note: For details, see section 4.



sensors. For example, the 700/670 nm ratio algorithm can reliably detect Chl-*a* at high concentrations  $>30 \text{ mg m}^{-3}$  and could be used to detect the presence of hypertrophic systems. Similarly, an NIR ratio algorithm could within reasonable error be used to detect high concentrations of TSS  $>30 \text{ g m}^{-3}$ , enabling sediment plume monitoring in estuaries. Water clarity ( $z_{SD}$ ) can be determined using a red/blue ratio; the suggested band ratios could be used to detect PC-containing cyanobacterial blooms, albeit with a lesser degree of confidence and depending on the availability of suitable sensors, to provide warning information to lake managers. Therefore, there are specific applications, for example, eutrophication assessment, sediment plume monitoring, water clarity assessments, for which the empirical approach has a great deal to offer.

A certain degree of error will inevitably be associated with any estimates from remote sensing; nevertheless, an acceptable level of error should be determined using sensitivity analyses (e.g. Hu, 2009) or *in situ* validation. The size of the error will depend on the magnitude of the signal from the parameter (large = less error, small = greater error), the instrumental design (hyperspectral/broad band, SNR), the data quality (e.g. atmospheric correction if used) and the degree to which the algorithm is parameterized for the system under investigation. With current instrumental constraints, and allowing for the complexity of detection, it is important to recognize that certain parameters such as  $a_{CDOM}$  and PC can be estimated with less confidence than others. Targets with smaller signals, such as clear-water oligotrophic systems, will also remain more challenging than systems with larger signals, for example, hypertrophic or sediment-laden waters. The error associated with estimates means that, in some instances, the information could be viewed as qualitative rather than explicitly quantitative, although quantitatively accurate estimates are desirable. Thus, although accepting a certain degree of error that remains to be determined, a great amount of useful information is readily accessible using the empirical approach.

Global assessments of the quality, quantity and changes occurring in inland fresh and transitional waters from remote sensing are likely to become commonplace within the next few decades. Remote sensing will play a vital role in determining the growing impact that global change will have on these limited and increasingly valuable resources. The achievement of this goal will depend on the availability of suitable techniques capable of providing reliable information on a regional and possibly even larger scale. This might be realized through moving towards the development not only of model-based algorithms but also of empirically based algorithms for common use in inland and transitional waters. The empirical approach, which is easy to implement and computationally simple, could play an important role alongside more advanced modelling procedures in the future of water remote sensing, especially for operational near real-time water-observation systems. As satellite sensors become more advanced in terms of spectral resolution, sensitivity and frequency of acquisition (even geostationary orbits), the detection of various parameters should improve, providing a wealth of information for water-quality monitoring and assessment. A number of outstanding issues still need to be addressed with regard to empirical algorithms. There is a need for cooperative parameterization and validation of empirical algorithms facilitated by a collective database containing *in situ* measurements in inland and transitional waters. Error assessment and sensitivity analyses should also be performed as part of this process to gain estimates of the accuracy and precision of the algorithms. Finally, large quantities of archived data from sensors such as MODIS and MERIS containing a wealth of potential information on inland and transitional waters have yet to be processed.

This article has provided a review of current empirical methods of retrieving various biogeophysical and optical parameters from remotely sensed data in inland and transitional waters. It provides an overview of the optimal current empirical procedures for estimating these parameters in a diversity of inland and coastal waters from a variety of remotely sensed instruments mounted on *in situ*, airborne and satellite platforms. Importantly, the review has not included more advanced inverse radiative transfer modelling or inversion algorithms, and readers are encouraged to find these elsewhere (e.g. IOCCG 2000, 2006). The review demonstrates the considerable number of recent studies using empirical procedures and the continual rapidly growing interest in water remote sensing. Therefore, although trends in bio-optical remote sensing point towards the use of more advanced inversion type models based on radiative transfer, which offer a suite of simultaneously derived optical and biogeophysical products (e.g. Odermatt *et al.* 2008), the usefulness of simpler yet robust empirical procedures is palpable. In operational monitoring contexts, the empirical approach has a demonstrable capability to provide timely and accurate information for a variety of parameters in lakes and estuaries that can be used for a diversity of applications. Current and future space-borne remote-sensing instruments have also been reviewed. It is hoped that the review will provide a useful reference for workers seeking to employ empirical procedures of remote sensing in inland and transitional Case 2 waters.

### Acknowledgements

I would like to express my sincere thanks to the University of Cape Town postgraduate publication incentive scholarship, one anonymous reviewer for their useful comments and criticisms, and my supervisor Dr. Stewart Bernard for encouraging me to write this article.

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**Appendix A: List of abbreviations in table 1****Data Range**

FNU = Formazin nephelometric units

*Data type*

AISA = Airborne imaging spectrometer for applications

ALI = Advanced land imager

AMMS = Airborne multispectral measurement system

ASTER = Advances space-borne thermal emission and reflection radiometer

CASI = Compact airborne spectrographic imager

CHRIS = Compact high-resolution imaging spectrometer

LISS III = Linear imaging self-scanning sensor

LS 7 ETM+ = Landsat 7 enhanced thematic mapper

LS TM = Landsat thematic mapper

MERIS = Medium-resolution imaging spectrometer

MIVIS = Multispectral infrared and visible imaging spectrometer

MODIS = Moderate-resolution imaging spectrometer

ROSIS = Reflective optics system imaging spectrometer

SAR = Synthetic aperture radar

Sea WiFS = Sea-viewing wide field-of-view sensor

SEVIRI MSG = Spinning enhanced visible and infrared imager on

METEOSAT second generation

Spec. = Spectroradiometer

SPOT HRV = Le Systeme Pour l'Observation de la Terra High Resolution Visible

*Water quality parameters (WQPs)*

$a_{\text{CDOM}}$  = absorption by coloured dissolved organic matter

Chl-*a* = Chlorophyll-*a*

COST = Cosine transmittance

PC = Phycocyanin pigment

$z_{\text{SD}}$  = Secchi disk depth

SPIM = Suspended particulate inorganic material

TSS = Total suspended solids

Turb = Turbidity

*Atmospheric correction*

6S = Second Simulation of the Satellite Signal in the Solar Spectrum (Vermote *et al.* 1997)

DDV = Dense dark vegetation

DOS = Dark object subtraction

ELM = Empirical line method (Moran *et al.* 2001)

ATCOR = Atmospheric correction

LOWTRAN = Low-resolution atmospheric transmittance code

MODTRAN = Moderate-resolution atmospheric transmittance code

MUMM = Management unit of the North Sea mathematical models

RTC = Radiative transfer code

SeaDAS = SeaWiFS data analysis system

SMAC = Simplified method of atmospheric correction (Rahman and Dedieu 1994)

SOS = Successive order of scattering code (Lenoble *et al.* 2007)

*Statistical technique*

ANN = Artificial neural network

BOM = Bio-optical model

CHROM = Chromaticity analysis  
GEGA = Grammatical evolution genetic algorithm  
LMM = Linear mixture modelling  
LR = Linear regression  
LSU = Linear spectral unmixing  
LT-LR = Log-transformed linear regression  
LT-MLR = Log-transformed multiple linear regression  
LT-MLSR = Log-transformed multiple linear stepwise regression  
MIP = Modular inversion and processing system  
MLR = Multiple linear regression  
NLR = Non-linear regression  
NN = Neural network  
Poly = Polynomial regression (number indicated order)  
PCA = Principal component analysis  
SA = Semi-analytical  
SAM = Spectral angle mapper  
SDA = Spectral decomposition algorithm