Table 2
The rate of oxygen consumption by various cells in culture.

Cell line or tissue	Cell type	OCR (amol cell ⁻¹ s ⁻¹)	OCR, original units (as reported)	Comments, methods (cell growth conditions)	Ref.
HL-60	Human promyelocytic leukemia (SC)	7.5	0.40-0.50 fmol min ⁻¹ cell ⁻¹	Fick's law (G1) ^a	[71]
HL-60	Human promyelocytic leukemia (SC)	11.5	11.46 ± 0.40^{f} pmol $O_2 s^{-1} (10^6 cells)^{-1}$	Oxygen monitor with Clark electrode (G1) ^a	[72]
HL - $60p^0$	Leukemia cells with knockout mitochondria (SC)	4.7	$4.74 \pm 0.16^{\rm f}$ pmol O_2 s ⁻¹ $(10^6$ cells) ⁻¹	Oxygen monitor with Clark electrode (G1) ^a	[72]
U937	Human histocytic leukemia (SC)	5.0	0.30 fmol min ⁻¹ cell ⁻¹	Fick's law (G1) ^a	[71]
U937	Human histocytic leukemia (SC)	11.0	11.00 ± 0.83^{f} pmol $O_2 s^{-1} (10^6 cells)^{-1}$	Oxygen monitor with Clark electrode (G1) ^a	[72]
Jurkat	Human acute lymphoblastic leukemia (SC)	12	$11.89 \pm 0.50 \; pmol \; O_2 \; s^{-1} \; (10^6 \; cells)^{-1}$	Oxygen monitor with Clark electrode (G1) ^a	[72]
MDCK	Dog kidney (AC)	20.8	1.25 fmol min ⁻¹ cell ⁻¹	Fick's law (G1) ^a	[71]
WEHI	Murine myelomonocytic leukemia cell line (SC)	7	0.4 fmol min ⁻¹ cell ⁻¹	Fick's law (G1) ^a	[71]
WEHI213	Murine myelomonocytic leukemia cell line	9.4	$9.44 \pm 0.48^{\rm f}$ pmol O_2 s ⁻¹ $(10^6$ cells) ⁻¹	Clark electrode	[72]
MCL5	Lymphoblastoid (SC)	3.5	0.21 fmol min ⁻¹ cell ⁻¹	Fick's law	[71]
CH2	Lymphoblastoid (SC)	5.8	0.35 fmol min ⁻¹ cell ⁻¹	(G1) ^a Fick's law (G1) ^a	[71]
Ehrlich ascites tumor cells	Mouse carcinoma (SC)	27	27 amol cell ⁻¹ s ⁻¹	Warburg apparatus	[43,45]
ALMA-16	Hybridoma (SC)	13	0.8 fmol min ⁻¹ cell ⁻¹	Fick's law (G1) ^a	[71]
Hybridoma	Murine hybridoma (SC)	61	0.22 pmol cell ⁻¹ h ⁻¹	Respirometer	[73]
C6	Rat glial tumor (on Cytodex beads)	12	0.7 fmol min ⁻¹ cell ⁻¹	Fick's law (G1) ^a	[71]
C6	Rat glial tumor (SC)	12	0.7 fmol min ⁻¹ cell ⁻¹	Fick's law (G1) ^a	[71]
WI-38	Human embryonic lung fibroblasts (on Cytodex beads)	2.5	0.15 fmol min ⁻¹ cell ⁻¹	Fick's law (G1) ^a	[71]
WI-38	Human embryonic lung fibroblasts (on Cytodex beads)	1.7	0.10 fmol min ⁻¹ cell ⁻¹	Fick's law	[71]
A20	Mature murine B cell lymphoma (SC)	10	$9.67 \pm 0.50^{\rm f} \ {\rm pmol} \ {\rm O_2 \ s^{-1}} \ (10^6 \ {\rm cells})^{-1}$	Clark electrode (G1) ^a	[72]
EL4	Murine T cell lymphomas (SC)	7.7	7.69 ± 0.40^f pmol O_2 s $^{-1}$ $(10^6$ cells $)^{-1}$	Clark electrode (G1) ^a	[72]
P815	Murine mastocytoma cell line (SC)	5.2	$5.15 \pm 0.37^{\rm f}~{ m pmol}~{ m O}_2~{ m s}^{-1}~(10^6~{ m cells})^{-1}$	Clark electrode (G1) ^a	[72]
BW1100	Murine mastocytoma cell line (SC)	8.1	$8.11 \pm 0.35^{\rm f} \ {\rm pmol} \ {\rm O_2 \ s^{-1}} \ (10^6 \ {\rm cells})^{-1}$	Clark electrode (G1) ^a	[72]
D2SC/1	Murine dendritic cell line (SC)	12.6	$12.56 \pm 0.83^{\rm f}$ pmol O ₂ s ⁻¹ $(10^6 \text{ cells})^{-1}$	Clark electrode (G1) ^a	[72]
MEF	Mouse embryonic fibroblasts	7	0.4 nmol min ⁻¹ (10 ⁶ cells) ⁻¹	Seahorse XF24 analyzer	[74]
MEF	Mouse embryonic fibroblasts	60	3.6 fmol min ⁻¹ cell ⁻¹	Seahorse XF24 analyzer	[75]
Myocytes	Neonatal cardiomyocytes	100	300 pmol min ⁻¹ (50,000 cells) ⁻¹	Seahorse XF24 analyzer	[76]
NRVM primary cell culture TIME cells	Neonatal rat ventricular myocyte (AC) Tert-immortalized microvascular endothelial cells	40 28	180 pmol min ⁻¹ (75,000 cells) ⁻¹ 50 pmol min ⁻¹ (30,000 cells) ⁻¹	Seahorse XF24 analyzer Seahorse XF24 analyzer	[52] [77]
Podocytes	Primary mouse podocytes (a kidney epithelial cell)	83	100 pmol min ⁻¹ (20,000 cells) ⁻¹	Seahorse XF24 analyzer	[78]
MC3T3 (on polysaccharide scaffolds)	Mouse myoblast (AC)	13	0.80 fmol min ⁻¹ cell ⁻¹	Fick's law (G1) ^a	[71]
C2C12	Mouse myoblast (on HA-FN scaffold)	3.7	0.22 fmol min ⁻¹ cell ⁻¹	Fick's law	[71]
Rat fibroblasts	Rat 1a spontaneously immortalized rat embryo fibroblasts	190	225 pmol min ⁻¹ (20,000 cells) ⁻¹	Seahorse XF24 analyzer	[79]

(continued on next page)

Cell line or tissue	Cell type	OCR (amol cell ⁻¹ s ⁻¹)	OCR, original units (as reported)	Comments, methods (cell growth conditions)	Ref.
Rat hepatocytes (fresh)	Primary, rat (SC)	200	12 fmol min ⁻¹ cell ⁻¹	Fick's law	[71]
Rat hepatocytes (fresh)	Primary, rat (on scaffold)	200	12 fmol min ⁻¹ cell ⁻¹	Fick's law	[71]
Rat hepatocytes	Rat hepatocytes	350	0.35 nmol s ⁻¹ (10 ⁶ cells) ⁻¹	Clark electrode with real-time numerical averaging	[49]
Rat hepatocytes	Rat hepatocytes	430	0.43 nmol s ⁻¹ (10 ⁶ cells) ⁻¹	Clark electrode	[51]
Porcine hepatocytes	Day 4 after seeding	900	0.9 nmol s ⁻¹ (10 ⁶ cells) ⁻¹	Clark electrode with real-time	[49]
rotelle nepatocytes	Day 15 after seeding	300	0.3 nmol s ⁻¹ (10 ⁶ cells) ⁻¹	numerical averaging	[49]
C	Rat brain, no treatment	(65 amol s ⁻¹	3.92 nmol min ⁻¹ (mg protein) ⁻¹	Clark electrode	1001
Synaptosomes	kat brain, no treatment	ng-protein-1)			[80]
Sf9 insect cells	Spodoptera frugiperda, ovarian	33	2.0 fmol min ⁻¹ cell ⁻¹	Fick's law (G2) ^b	[71]
Hi-5 insect cells	Trichoplusia ni, ovarian	105	6.3 fmol min ⁻¹ cell ⁻¹	Fick's law	[71]
FS-4	Human diploid foreskin cells (SC)	14	0.05 mmol h ⁻¹ (10 ⁹ cells) ⁻¹	(G2) ^b Based on oxygen demand by cells and mass transfer coefficient (G3) ^c	[48]
HLM	Liver (AC)	102	0.37 mmol h ⁻¹ (10 ⁹ cells) ⁻¹	Use modified Cartesian diver	[48,81]
LIR	Liver (AC)	83	0.30 mmol h ⁻¹ (10 ⁹ cells) ⁻¹	Use modified Cartesian diver	[48,81]
Skin fibroblast	Human (AC)	18	0.064 mmol h ⁻¹ (10 ⁹ cells) ⁻¹	Use modified Cartesian diver	[48,81]
143B	Human osteosarcoma (AC)	16.3	16.32 ± 0.53f pmol O ₂ s ⁻¹ (10 ⁶ cells) ⁻¹	Oxygen monitor with Clark electrode	[72]
143Bp0	Human osteosarcoma with knockout	5.6	$5.62 \pm 0.40^{\text{f}} \text{pmol O}_2 \text{ s}^{-1} (10^6 \text{ cells})^{-1}$	Oxygen monitor with Clark electrode	[72]
ТЧЭБРО	mitochondria (AC)	5.0	,	Oxygen monitor with clark electrode	
Detroit 6	From bone marrow of lung cancer patients (AC)	120	0.43 mmol h ⁻¹ (10 ⁹ cells) ⁻¹		[82]
MCN	Leukemia (AC)	61	0.22 mmol h ⁻¹ (10 ⁹ cells) ⁻¹	Based on oxygen demand by cells and mass transfer coefficient	[82]
Conjunctiva	Human eye cells (AC)	78	0.28 mmol h ⁻¹ (10 ⁹ cells) ⁻¹	Based on oxygen demand by cells and mass transfer coefficient	[82]
Lung To	Human embryonic lung cells (AC)	67	$0.24 \text{mmol h}^{-1} (10^9 \text{cells})^{-1}$	Based on oxygen demand by cells and mass transfer coefficient	[82]
Intestine 407	Human (AC)	111	0.40 mmol h ⁻¹ (10 ⁹ cells) ⁻¹	Based on oxygen demand by cells and mass transfer coefficient	[82]
MAF-E	Adult Fallopian tube (AC)	106	0.38 mmol h ⁻¹ (10 ⁹ cells) ⁻¹	Based on oxygen demand by cells and mass transfer coefficient	[82]
Red blood cells	Human (adult)	4×10 ⁻⁵	Contribution estimated from the rate of autoxidation of oxyhemoglobin to form superoxide; H ₂ O ₂ is generated at a rate of 3.9 ± 0.6 nmol·h ⁻¹ ·gHb ⁻¹	This corresponds to about 50 superoxide radicals being produced each second in an RBC.	[83]
Red blood cells	Rabbit	0.02	$(1.5 \pm 0.2) \times 10^{-15} \text{ L RBC}^{-1} \text{ h}^{-1}$	Gilson differential recording respirometer, 38 °C	[84]
Lymphoblastoid (Namalioa)	Human (AC)	15	0.053 mmol h ⁻¹ (10 ⁹ cells) ⁻¹	Based on oxygen demand by cells and mass transfer coefficient	[85]
J774A.1	Murine macrophages (AC)	31	1.87 nmol min ⁻¹ (10 ⁶ cells) ⁻¹	EPR oximetry	[86]
J774A.1	Murine macrophages (AC)	6.2	6.18 ± 0.33 ^f pmol O ₂ s ⁻¹ (10 ⁶ cells) ⁻¹	Oxygen monitor with Clark electrode	[72]
CHO	Chinese hamster ovary cells (SC)	74	4.43 nmol min ⁻¹ (10 ⁶ cells) ⁻¹	EPR oximetry	[86]
CHO	Cliniese nanister ovary cens (SC)	74	4.45 IIII0I IIIII (10 Cells)	(G4) ^d	[00]
CHO	Chinese hamster ovary cells (SC)	88	3.2×10 ⁻¹³ mol cell ⁻¹ h ⁻¹ (5.3 nmol min ⁻¹ (10 ⁶ cells) ⁻¹)	Microtiter plate with oxygen sensor	[87]
CHO	Chinese hamster ovary cells (SC)	86	0.31 pmol cell ⁻¹ h ⁻¹	Using a respirometer	[73]
CHO	Chinese hamster ovary cells (SC)	8.0	0.50 fmol min ⁻¹ cell ⁻¹	Fick's law (G1) ^a	[71]
CHO	Chinese hamster ovary cells (SC)	63	3.8×107 molecules of O2 s-1 cell-1	EPR oximetry	[47]
CCD	Kidney cortex collecting duct cells	25	1.48 nmol min ⁻¹ (10 ⁶ cells) ⁻¹	EPR oximetry	[86]
AG08472	Vascular endothelial cells of the pig	17	1±0.15 nmol min ⁻¹ (10 ⁶ cells) ⁻¹	Optical method using oxygen quenchers	[88]
1100-112	thoracic aorta (AC)	.,	(when measured at 22 °C), 0.64 (at 4 °C)	operati method using oxygen quenchers	[oo]
AG08473	SMC of cells of the pig thoracic aorta (AC)	44	2.64 ± 0.14 nmol min ⁻¹ (10 ⁶ cells) ⁻¹	Optical method using oxygen quenchers	[88]

Table 2 (continued)

Cell line or tissue	Cell type	OCR (amol cell-1 s-1)	OCR, original units (as reported)	Comments, methods (cell growth conditions)	Ref.
HeLa cells	Human cervical carcinoma cells (AC)	(200 amol s ⁻¹ ng-protein ⁻¹)	$11.7\pm1.3~\mathrm{nmol~min^{-1}}~\mathrm{(mg~protein)^{-1}}$	Clark electrode (G1) ^a	[89]
HeLa cells	Human cervical carcinoma (AC)	12.5	$12.50 \pm 0.5^{\rm f}$ pmol O ₂ s ⁻¹ ($10^{\rm 6}$ cells) ⁻¹	Clark electrode (G1) ^a	[72]
A549	Human adenocarcinoma alveolar epithelial	27	1.6 nmol min ⁻¹ (10 ⁶ cells) ⁻¹	Seahorse XF24 analyzer	[90]
NIH-H460	Human large cell lung cancer, epithelial	30	1.8 nmol min ⁻¹ (10 ⁶ cells) ⁻¹	Seahorse XF24 analyzer	[90]
L-6 myoblasts	Human muscle (AC)	(200 amol s ⁻¹ ng-protein ⁻¹)	12 ± 1.3 nmol min ⁻¹ (mg protein) ⁻¹	Clark electrode (G1) ^a	[89]
Beating cardiac myocytes	New born rats	(680 amol s ⁻¹ ng-protein ⁻¹)	40.5 ± 1.3 nmol min ⁻¹ (mg protein) ⁻¹	Clark electrode with Lucite attachment (G1) ^a	[89]
Beating cardiac myocytes	Old rats	(1200 amol s ⁻¹ ng-protein ⁻¹)	69.5 nmol min ⁻¹ (mg protein) ⁻¹	Clark electrode with Lucite attachment (G1) ^a	[91]
Heart nonmuscle	Newborn rat	(200 amol s ⁻¹ ng-protein ⁻¹)	$11.8 \pm 0.7 \text{ nmol min}^{-1} \text{ (mg protein)}^{-1}$	Clark electrode (G1) ^a	[89]
Bovine endothelial	From aortae of cattle (AC)	(67 amol s ⁻¹ ng-protein ⁻¹)	$4.0\pm0.7~\mathrm{nmol~min^{-1}}~\mathrm{(mg~protein)^{-1}}$	Clark electrode (G1) ^a	[92]
Renal mesangial	Rat cells (AC)	(150 amol s ⁻¹ ng-protein ⁻¹)	$9.0\pm0.3~\mathrm{nmol~min^{-1}}~\mathrm{(mg~protein)^{-1}}$	Clark electrode (G1) ^a	[92]
LLC-PK	Renal epithelial cells from pig kidney (AC)	(320 amol s ⁻¹ ng-protein ⁻¹)	$19.0 \pm 0.9 \mathrm{nmol min^{-1}} (\mathrm{mg protein})^{-1}$	Clark electrode (G1) ^a	[92]
LLC-MK	Rhesus monkey kidney (AC)	(470 amol s ⁻¹ ng-protein ⁻¹)	28.2 ± 0.7 nmol min $^{-1}$ (mg protein) $^{-1}$	Clark electrode (G1) ^a	[92]
HepG2	Human hepatoma cells (AC)	(110 amol s ⁻¹ ng-protein ⁻¹)	$6.7\pm1.2~\mathrm{nmol~min^{-1}}~\mathrm{(mg~protein)^{-1}}$	Clark electrode (G1) ^a	[92]
Hep3B	Human hepatoma cells (AC)	(160 amol s ⁻¹ ng-protein ⁻¹)	$9.6\pm1.4~\mathrm{nmol~min^{-1}}~\mathrm{(mg~protein)^{-1}}$	Clark electrode (G1) ^a	[92]
AFP-27	Murine hybridoma cell line	6.0	2.15×10 ⁻⁸ μmol cell ⁻¹ h ⁻¹	Tissue oxygen probe system, 37 °C (G1) ^a	[93]
Human mesenchymal preadipocytes	Undifferentiated (AC)	25	$0.591 \pm 0.302 \text{ nmol min}^{-1} (0.4 \times 10^6 \text{ cells})^{-1}$	Clark electrode (G1) ^a	[94]
Human mesenchymal preadipocytes	Differentiated (AC)	120	$2.865 \pm 0.219 \text{ nmol min}^{-1} (0.4 \times 10^6 \text{ cells})^{-1}$	Clark electrode (G1) ^a	[94]
RAW264.7	Transformed mouse macrophages (AC)	8.9	$8.89 \pm 0.23^{\rm f}$ pmol O_2 s ⁻¹ (10^6 cells) ⁻¹	Clark electrode (G1) ^a	[72]
ВНК	Baby hamster kidney	83	0.3 pmol cell ⁻¹ h ⁻¹	Respirometer (G5) ^e	[73]
TM4	Murine testicular cells	(10 amol s ⁻¹ ng-protein ⁻¹)	37 nmol h ⁻¹ (mg protein) ⁻¹	Polarography at 34 °C	[95]
MCF-7	Breast cancer cell line (AC)	(1300 amol s ⁻¹ ng-protein ⁻¹)	77.5 nmol min ⁻¹ (mg protein) ⁻¹	Clark electrode (G1) ^a	[96]

Table 2 (continued)

Cell line or tissue	Cell type	OCR (amol cell ⁻¹ s ⁻¹)	OCR, original units (as reported)	Comments, methods (cell growth conditions)	Ref.
Molt-4 cells	Human leukemia cell line (AC)	12	0.7 nmol min ⁻¹ (10 ⁶ cells) ⁻¹	EPR with 15N-PDT, 37 °C	[97]
				(G1) ^a	
Molt-4 ρ ⁰ cells	Human leukemia cell line with knockout mitochondria (AC)	1.3	0.08 nmol min ⁻¹ (10 ⁶ cells) ⁻¹	EPR with ¹⁵ N-PDT, 37 °C	[97]
LNCAP	Prostate cancer (AC)	63	$3.75 \pm 1.12 \text{ nmol min}^{-1} (10^6 \text{ cells})^{-1}$	EPR with 15N-PDT, 37 °C	[97]
AGS	Human gastric cancer cell line (AC)	27	1.6 nmol min ⁻¹ (10 ⁶ cells) ⁻¹	Clark electrode, 25 °C	[98]
BM MNCs	Human bone marrow mononuclear cells	10.6	0.038 (adherent) µmol h ⁻¹ (10 ⁶ cells) ⁻¹	Hermetically sealed tissue culture	[99]
	Cultured for 14 days	6.9	0.025 (nonadherent) μ mol h ⁻¹ (10 ⁶ cells) ⁻¹	well inserts equipped with oxygen	
				electrodes, 37 °C	
Escherichia coli	Bacteria	0.13	0.008 fmol min ⁻¹ cell ⁻¹	Fick's law	[71]
	Book 1		0.0016 1 1 =1 11=1	(G3)°	
Salmonella typhimurium	Bacteria	0.017	0.001 fmol min ⁻¹ cell ⁻¹	Fick's law (G3) ^c	[71]
Saccharomyces cerevisiae	Brewer's yeast (Edme)	2	0.12 fmol min ⁻¹ cell ⁻¹	Fick's law	[71]
	aremer's years (manne)			(G2) ^b	10.01
Candida albicans	Yeast (fungus)	25 (per cfu)	1.5 fmol min ⁻¹ cfu ⁻¹	Fick's law	[71]
				(G2) ^b	
Embryonic stem cells	Murine (AC)	40	4×10 ⁻¹⁷ mol s ⁻¹ cell ⁻¹	Using oxygen probe (Phoenix	[100]
				Electrode Co., Houston, TX, USA)	
Neural stem cells	Murine (AC)	31	3.06×10 ⁻¹⁷ mol s ⁻¹ cell ⁻¹	Using oxygen probe (Phoenix	[101]
				Electrode Co., Houston, TX, USA)	
Human, adult neutrophils	Preincubated with chemotactic factor	86	5.16 nmol min ⁻¹ (10 ⁶ neutrophils) ⁻¹	Clark electrode, 37 °C	[102]
	(FMLP) and activated with OPZ				
Human neutrophils	(opsonized zymosan) Polymorphonuclear neutrophils (PMN)	15	4.38 nmol min ⁻¹ (5×10 ⁷ neutrophils) ⁻¹	Clark electrode, 37 °C	[103]
Human neutrophils	PMN activated with LPS	16	4.87 nmol min ⁻¹ (5×10 ⁷ neutrophils) ⁻¹	Clark electrode, 37 °C	[103]
Human neutrophils	PMN when phagocytizing E. coli	16	48.6 nmol min ⁻¹ (5×10 ⁷ neutrophils) ⁻¹	Clark electrode, 37 °C	[103]
Human neutrophils	PMN when phagocytizing	34	102 nmol min ⁻¹ (5×10 ⁷ neutrophils) ⁻¹	Clark electrode, 37 °C	[103]
	Staphylococcus aureus		(onto headophila)		[105]
Human neutrophils	PMN when phagocytizing zymosan	25	73.9 nmol min ⁻¹ $(5 \times 10^7 \text{ neutrophils})^{-1}$	Clark electrode, 37 °C	[103]
Rat neutrophils	Untreated	8.3	299.3 nmol h ⁻¹ (10 ⁷ cells) ⁻¹	Clark electrode, 37 °C	[104]
Rat neutrophils	Treated with glucose (5 mM)	9.1	328.2 nmol h ⁻¹ (10 ⁷ cells) ⁻¹	Clark electrode, 37 °C	[104]

AC, adherent cells; cfu, colony-forming units; EPR, electron paramagnetic resonance; OCR, rate of oxygen consumption; SC, suspension cells; ¹⁵N-PDT, 4-oxo-2, 2, 6, 6-tetramethylpiperidine-d¹⁶-1-¹⁵N-oxyl.

a C1, cells grown at 37 °C, with 5% C0, 95% humidity.

b C2, cells grown at 27 °C, in a humidity chamber.

c G3, cells grown at 37 °C.

d G4, cells grown in spinner flasks, 37 °C, 12% CO₂, 88% humidity.

d G4, cells grown in spinner flasks, 37 °C, 12% CO₂, 88% humidity.

e G5, cells grown at 35 °C.

f Contributions from cell surface, basal, and mitochondrial O₂ consumption are given in Table 4,