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KEY WORDS

life-testing, electric vehicles, lead-acid batteries, battery chargers, gas pressure control

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A SYSTEM EVALUATION OF LEAD-ACID BATTERY CHARGERS

J L Weininger and E.G. Siwek

INTRODUCTION

In this study, three lead-acid battery charge control methods are evaluated Evaluation is made on the basis of battery cycle life, changes in charging efficiency during successive cycling, and the effects of battery design on charger/battery interaction.

Lead-acid battery charging systems which use gases evolved from the battery during charge as a means for charging control are potentially more efficient than conventional battery charging systems. Further, it is expected that gas-controlled charging systems can result in prolonged life because they avoid elevated temperatures associated with overcharging and excessive gassing which loosens active materials from the plate structure during overcharge.

Below, the selection of a suitable current profile is discussed, followed by a description of the three charging methods. Details are given for the two types of batteries tested, for the testing procedure, and the resulting battery performance throughout cycle life. The report is concluded with a discussion of the results

CURRENT PROFILE SELECTION

For this program, maximum charging efficiency was desired as well as maximum battery life. These two goals are consistent since an efficient charge, because of its minimal gassing and minimal temperature rise during charge, results in prolonged battery life.

The charging current-time profile which results in a charge of maximum efficiency is that current profile which follows or coincides with the battery charge acceptance curve. This is illustrated by the idealized charge acceptance curve of Fig 1. The curve corresponds to a max mum acceptable initial charging current and a finishing current just sufficient to maintain the battery above its self-discharge rate. In the case of one of the 12 amp-hr batteries tested (battery A below), these ideal values previously had been found to be 55 and 0.12 amp. However, for practical purposes, because initial extremely high charging currents were too difficult to manage within the scope of this work, a 2C initial charging rate (24 amp) was used for the three charging modes under study. Similarly, the end of charge rate proved to be too small especially for aging batteries, so that a finishing rate of 0.24 to 0 48 amp (C/50 to C/24) was chosen in this work.

CHARGING METHODS

Of the three selected charging modes, two were based on gas control, by either mass flow or differential pressure control. The third method was a

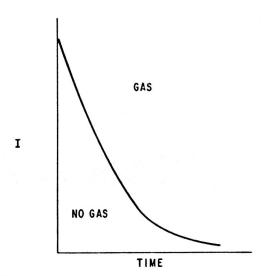


Fig. 1 Idealized charge acceptance curve.

Modified Constant Potential Method (MCP), which was chosen for comparison as a more conventional charging method. All three of these charging methods rely upon an exponentially decaying charging current, capable of reflecting and responding to the battery's diminishing charge acceptance during charging

The conventional MCP charging method is the simplest of the three charging modes selected. For this type of charge, two parameters are selected: the initial charging current and the constant charging voltage. The term "modified," when applied to this charging method, indicates that the initial charging current has been limited so that large current spikes are avoided at the beginning of charge, as would be the case at the beginning of charge of a fully discharged battery

Although the MCP charging method is the most popular of the conventional lead-acid battery charging systems, it has several inherent disadvantages. Some of these disadvantages result from the fact that, in some lead-acid batteries, the gassing potential is lowered with battery age. Since it is not convenient to adjust the charge bus voltage to compensate for this change, aging batteries gas more profusely and progressively earlier in the charge. This results in not only less efficient charging but in shortened battery life.

Another disadvantage results from the manner in which the charging current is regulated by this charging method. The battery voltage approaches the charger bus voltage asymptotically. The result is that, as the battery approaches full charge, more time is required to apply progressively less charge to the bat tery. A 100 percent charge is virtually impossible by this method, or for any charging method which em

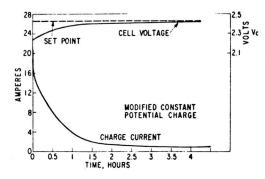


Fig 2 Charging current and voltage profile, modified constant potential charge method.

oys an exponentially decaying charging current profile (see Fig. 2).

The Mass Flow (FOMF) controlled charge is characterized by a charging current which is inversely proportional to the flow rate of gas from either or both electrodes of a charging battery At the beginning of charge, when no gas is being evolved from either electrode, the charging current is at its preset maximum, 24 amp, for this study As gas is evolved, the charging current is diminished so that the gas flow rate does not exceed the set point. Since the charge acceptance of the battery diminishes with charge, progressively less charging current is required to maintain the preset gas flow Finally, at the end of charge all of the charging current results in gassing. By definition, the battery is 100 percent charged and being maintained in the charged state by an overcharge of 0.24 amp, all of which results in the electrolysis of water (see Fig. 3). Based on a theoretical 100 percent gassing of 11 2 cc min/cell/amp, this corresponds to a gas flow of 11.2 × 0.24 = 2.688 cc min/cell.

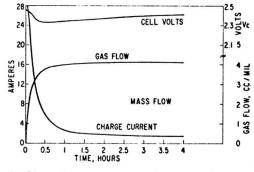


Fig. 3 Charging current, voltage, and mass flow profiles; first-order mass flow charge

While this charging method is not controlled directly by a voltage difference as in the case of the MCP method, and is, therefore, not immediately sensitive to battery age, it does require the battery to be sealed. Further, it requires a gas flowmeter with a voltage output, and some means for employing this voltage output in controlling the output of a charging power supply A tapering charging current, sim-

ilar to that realized in the MCP charging, also results in progressively longer times to charge active electrode material as the battery approaches full charge. This is inevitable where extremely high-efficiency charging is desired by a charging method which utilizes a tapering current profile which approaches exponential form.

The Gas Pressure (SOGP) controlled charge is characterized by a charging current that decays in direct ratio to the amount of evolved gas necessary to maintain a preselected differential gas pressure within the battery container Since this system allows gas to escape from the battery container via a small controlled orifice, it is also controlled by gas flow. Unlike the FOMF charge control method, control does not begin until the preselected differential pressure is reached. This results in a charging current profile which begins as a constant current of 24 amp and remains at 24 amp until the set point 0 4 psig is reached. At this point, the charging current begins * to diminish such that the differential pressure is maintained. This calibrated differential pressure corresponds to the same flow of gas from the battery as for the FOMF-controlled charging method. Shown in Fig. 4 are the current, voltage, and pressure profiles used for this charging method.

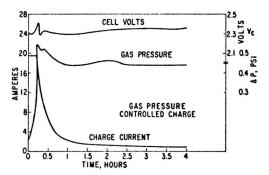


Fig. 4 Charging current, voltage, and differential pressure profiles; second-order gas pressure charge method.

This method requires a pressure transducer with a voltage readout and a means for using that readout to control the output of the charging circuit power supply.

Since both gas-controlled charge methods described in this report require sealed batteries, they present potential hazards resulting from runaway charging. In practice, thermal overload or overvoltage devices can be utilized in the prevention of runaway during charging. Life-testing control circuits, used for this program, included overvoltage control devices which worked well. Other potential problems associated with this type of charge control can be solved with present state-of-the-art electronic and other controls.

DESCRIPTION OF BATTERIES

All of the data for this program were obtained on two types of lead-acid batteries, both of which have cast lead-antimony grids. Each of the battery types contained three cells and was rated by its manufacturer as 12 amp-hr. The chief differences between the two battery types were the number of plates per cell and the discharge rates used in rating the batteries. Also, grids of Type A had a lower antimony content (4%) than Type B grids (6% Sb).*

EXPERIMENTAL PROCEDURE

Data Logging

Battery characterization consisted of carefully monitoring the following seven parameters during charging:

Time	Hr
Battery voltage	Vb
Positive electrode to ${\rm Hg/Hg_2SO_4}$	V+
Negative electrode to $\mathrm{Hg}/\mathrm{Hg}_2\mathrm{SO}_4$	V-
Charging current (amp)	Ιt
Gas flow (cc/min)	$\mathbf{F}_{\mathbf{t}}$
Oxygen in gas (%)	O_2
Temperature (°C)	T

The following were then calculated from the measured data:

Cell voltage	Vc
Oxygen flow (cc/min)	Fo
Hydrogen flow (cc/min)	Fh
Charging current, positive electrode	I+
Charging current, negative electrode	I-
Total amp-hour charge	AHt
AH charge, positive electrode	AH+
AH charge, negative electrode	AH-
Total watt-hour charge	WHt
WH charge, positive electrode	WH+
WH charge, negative electrode	WH-

From these computer calculated data, the following performance indicators were obtained from the batteries' discharge capacity (AH dis):

Total amp-hour efficiency (%) %AH (AH dis AHt)

Amp-hour efficiency, %AH+ (AH+/AHt)
positive electrode (%)

Amp-hour efficiency, negative electrode (%)	%AH- (AH-/AHt)
Total watt-hour efficiency (%)	%WH (WH dis/WHt)
Watt-hour efficiency, positive electrode (%)	%WH+ (WH+/WHt)
Watt-hour efficiency, negative electrode (%)	%WH- (WH-/WHt)
Positive charge retention (%)	%CR+ (AH dis AH+)
Negative charge retention (%)	%CR- (AH dis/AH-)

The above data logging procedure was followed while each of the batteries was being charged at constant current, 1 2 amp, and while each of the batteries was being charged under one of the three charging modes under investigation.

Experimental Data

Three constant current charge characterizations and three characterizations under one of the charging modes being investigated were run for each battery during its life as measured by the life-test cycling. Shown in Table I are the sequential steps followed by characterizing and life-testing of each of the six batteries.

The sequence of data gathering for the preconditioned batteries covered in this report was:

- Characterization using a C/10 charging rate for 12 hours followed by a C/10 discharge to 1.75 volts cell.
- Characterization using one of the three charging modes under investigation followed by a C/10 discharge to 1 75 volts cell.
- Repetitive charge-discharge cycling (lifetesting) using one of the charging methods under investigation and discharges at the C/10 discharge rate to 1.75 volts/cell.

In addition to characterization measurement of new batteries, characterization measurements were repeated when the battery under investigation had degraded to 75 percent of its initial discharge capacity Finally when the battery had degraded to 50 percent of its initial discharge capacity as determined from the life-testing portion of the program. For this program, a battery which had degraded to 50 percent of its initial stabilized discharge capacity was considered spent.

Six batteries were evaluated for this program: three Type A batteries and three Type B batteries. For purposes of identification, a code was used to identify charging mode, battery type, type of characterization, and characterization number. The code consists of three groups of letters followed by a digit, for example:

MCP-A-CC-1

^{*}For details of the battery construction and chemical analysis of the battery grids, see Final Report of ILZRO Project LE-205, Dec. 1975

[†]Assuming an average discharge voltage = 1.95 volts.

Table I

CHARACTERIZATION AND LIFE TESTING

Function	Cycle No.	Charge	Rate	Discharge	Rate	
Conditioning	1 to N	СС	C/10	CC	C/10	To develop stabil ze and establish buttery capacity
Characterization	N+1	cc	C/10	, a .	•	To characterize new battery at const I
Characterization	N+2	MCP	-	n		To characterize new battery at MCP
Life Test	(N+2) to M	MCP	-	n	"	Life Test to 75% of N
Characterization	M+1	cc	C/10	Ħ	XIII	Characterization at CC at 75% of N
Characterization	M+2	MCP	-	n ≬	n	Characterizat on at MCP at 75% of N
Tife Test	(M+2) to 1	MCP		on .	în .	Life Test to 50% of N
Characterization	L+1	сс	C/10	**	"	Characterization at CC at 50% of N
Characterization	L+2	MCP		: d =		Characterization at MCP at 50% of N

The first group of letters identifies this as a battery normally cycled at the MCP method. The second letter identifies the battery -- Type A. The third group of letters indicates the characterization method for a particular characterization: Constant Current (CC). The digit identifies a characterization number, (1). A all of six characterizations was planned for each cettery: three characterizations at constant current

and three characterizations at one of the charging modes. Codings for these six batteries are:

MCP-A MCP-B	Modified Constant Potential
FOMF-A FOMF-B	First-Order Mass Flow
SOGP-A	Second-Order Gas Pressure

As a distinction between the characterizations of the three different charging modes, Fig 5 presents the charge current as a function of time for the three methods.

COMPUTERIZED MANIPULATION OF CHARACTERIZATION DATA

From the above discussion of battery and characterization coding, it is apparent that in addition to the life-test data, 36 blocks of data were generated during the six characterization of six batteries.

In this report, only summary tables of the computer-generated data are shown. Table II shows the bottom lines of the computer readout sheets. Tables III and IV are summaries of the ampere-hour and watthour efficiencies as well as charge acceptance and discharge recovery calculated from the computer readout rheets.

Life-test data were obtained by continuous automatic cycling. In all cases, constant current discharges of 1 2 amp were terminated at 1 75 volts per cell. The actual cycle life-test data are given in Table V

DISCUSSION

Cycle Life

The longest-lived battery was MCP-B with 325 cycles. In this respect the MCP mode was best.

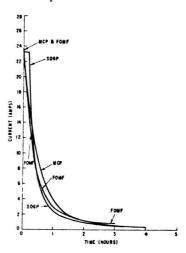


Fig. 5 Current profile for the three different charging modes.

†The complete listing of the 36 blocks of data is published in the Final Report of ILZRO Project No. LE-205, Dec. 1975, and can be obtained from the International Lead Zinc Research Organization, Inc., 292 Madison Avenue, New York, NY 10017

Table II END OF CHARGE CHARACTERIZATION DATA

			and Electronia		Current	Gas Flow	Per	• Gas	Flow	turi	rent				foral	Pos.	Neg.	
Experiment No.	Hrs.	Vc	Volts V+		Amps It	cc/min Ft	centage 0 ₂	Fo	FH	Ar I+	ips -	focal AH	AH+	tual #-	Energy W-Hr.	Elect W-Hr.	Elect. W-Hr.	AH Discharge
MCP-A-CC-I	12.00	2.73	37	31	1.20	. 69						14.40			13.76			12.00
MCP-A-CC-2	2.50 11.50	2.54	1.25	1.22	.20	.81 .69	33.7 36.1	3.92 0 97	1.72	0.15	0.15	14.94 14.12	10.84	1.52	34.39 31.25	24.25 27.84	25.87 28.94	12.50 8.60
MCP-MCP-I -2 ''	4.00 3.50 3.00 5.33 16.50 6.17	2.58 2.48 2.48 2.48 2.49 2.34	1.28 .27 1.28 1.30 1.34	1.30 1 21 .20 1 18 1.15 1.02	0.45 0.99 0.98 0./3 0.50 1.03	4.87 8.16 7.79 7.20 5.04 4.61	30.4 25.4 35.2 33.9 32.2	2.48 1.98 1.54 1.71 1.48	5.68 5.81 4.66 3.33 3.13	0.33 0.45 0.05 0.05 0.63	0.23 0.21 0.11 0.06 0.61	15.04 3.27 1.38 13.62 19.82 14.04	10.95 9.67 9.94 10.71 8.16	10.94 9.42 10.16 11.09 12.73	36.70 31.46 27.00 32.52 47.78 32.27	25./6 22.81 23.48 25.25 18.65	25.70 22.17 24.00 26.15 29.23	12.48 9.52 9.30 10.13 11.25 9.38
FOMF-A-CC-1	12.0	2.60	1.28	1.32	1.20	6.81	7.6	0.52	6.28	1.06	0.38	14.40	14.00	13.16	32.28	31.98	29.14	12.30
Battery Failure Battery Failure FOMF-A-FOMF-1	3.00	2.40	1.23	1.16	0.91	0.31	16.0	0.31	1.64	0.83	ი.69	12.34	11.93	11.90	29. 3	28.17	28.69	10.80
Battery Failure																		
Sogp-A-CC-1 -2 -3	12.50 12.33 16.00	2.47 2.51 2.51	. 27 . 34 . 34	1.30 1 17 1 17	1.25 1.28 1.25	1' 00 .25	28.7 40.9 33.8	4.30 4.60 4.44	10.7 6.65 4.31	0.10 0.05 0.05	0 0.39 0.07	15.04 15.54 19.07	12.47 8.36 10.39	12.76 9.85 10.77	33.08 36.73 44.65	26.88 19.01 23.26	27.47 22.62 24.10	10 14 10
SOGP-A-SOGP · I · 2 '' - 3	4.00 3.33 5.83	2.42 2.38 2.37	.25 .26 .25	1 18 1.12 12	0.44 0.76 0.49	3.88 3.37 2.75	17.7 26.1 27.4	0.69 0.89 0.75	3.19 2.48 2.00	0.26 0.52 0.29	0.01 0.43 0.32	12.90 7.09 11.74	10.45 5.99 10.30	10.46 6 17 10.48	30.95 15.89 27.04	25.09 13.31 23.70	25.10 13.72 24.10	6.30 10.60
MCP-8-CC-1 " -2 -3	12.17 12.00 8.50	2.70 2.45 2.34	1.31	. 38 1.12 1.08	1.20 1.22 1.24	10.50 7.74 4.25	29.8 35.3 39.7	3.13 2.73 .69	7.37 5.01 2.56	0.37 0.49 0.79	0.22 0.56 0.90	14.61 14.64 10.40	13.05 12.50 9.62	12.00 13.00 9.67	33.32 33.03 22.56	29.19 27.84 20.84	26.41 29.04 20.96	11 14 9.66 6.54
MCP-B-MCP-1 '' -2 '' -3	4.00 2.00 2.52	2.65 2.40 2.39	1.31 1.34 1.36	1.33 1.07 1.03	1.07 1.60 4.17	9.87 4.97 35.00	33.4 36.8 23.3	3.30 1.83 8.14	6.57 3.14 26.90	0.19 1 12 2.00	0.19 1.19 0.59	16.23 10.37 20.00	12.31 9.90 13.97	11.62 10.07 11.54	40.69 23.95 48.02	30.44 22.84 33.51	28.63 23.25 27.60	1.56 8.28 8.70
FOMF-B-CC-1 '' -2 '' -3	12.17 11.53 11.40	2.71 2.47 2.39	1.31 1.27 1.27	1.40 1.20 1.12	1.20 1.20 1.22	10.62 10.50 8.25	20.2 31.9 30.3	2,14 3.35 2.50	8.48 7.15 5.75	0.63 0.31 0.56	0.07 0.25 0.45	14.45 13.84 13.84	13.02 10.38 9.32	12.18 9.28 9.16	32.13 31.59 31.74	28.44 23.15 21.01	26.31 20.43 20.63	1.36 10.28 6.23
FOMF-8-FOMF-1 " -2 " -3	2.43 4.00 4.00	2.43 2.37 2.35	1.26 1.26 .28	1.17 1 11 1.07	1.08 0.42 0.61	1 17 3.00 3.50	30.5 30.9 32.2	0.36 0.43 1 13	0.82 2.07 2.37	0.99 0.17 0.31	0.98 0.14 0.29	13.16 9.49 6.83	12.91 8.44 5.73	13.04 8.46 5.74	31.42 22.48 16.20	30.81 20.01 13.62	31 11 20.06 13.04	10.56 8.92 5 40
SOGP-B-CC-1 -1* -2 3	23.00 (12.00) 11.50 11.50	2.48 (2.45) 2.36 2.37	.33 (1.26) '31 31	1 15 (1 18) 1.05 1.06	1 17 (1 18) 1.23 1.20	8.00 (5.62) 6.12 7.50	32.1 (32.6) 28.6 37.9	2.57 .83) 75 2.84	5.43 (3.79) 4.37 4.66	0.48 (0.69) 0.76 0.44	0.44 (0.67) 0.65 0.58	27.34 (14.42) 14.17 13.80	19.24 (13.44) 1 1 10.12	18.99 (13.40) 10.40 1 11	63.83 (31.88) 32.24 30.87	43.34 (29.50) 25.06 22.26	43.20 (29.40) 23.38 24.57	52 (11.25) 4.75 4.97
SOGP 8-SOGP-1 -2 -3	4.03 4.00 3.83	2.36 2.34 2.35	. 31 31	1.16	0.87 1.32 .42	2.50 6.75 6.87	24.0 36.6 31.5	0.60 2.47 2.16	1.90 4.29 4.71	0.71 0.66 0.84	0.62 0.75 1.79	13.20 11.89 10.39	9.36 8.14	9.64 9.64	30.94 27.66 24.56	23.5, 21 90 19.28	28.22 22 47 18.80	9.80 3.96 5.00

*Figure: In parentheses refer to charge ofter 12 hours not end of charge at 23 hours.

Table III SUMMARY DATA-CHARACTERISTICS OF BATTERIES WITH CONSTANT CURRENT MODE CHARACTERIZATIONS

Batt	Char		No of	Ampe	re Hou	r	Wat	t Hour		Percent Accept	Charge		very (%) arge***
No.	No.	Mode	Cycles	Charge	Dis.	Eff.	Charge	Dis.*	Eff.	Pos	Neg	Pos	Neg
MCP-A	1	CC	1	14 4	12 0	83	33 8	23 3	69	-	-	-	-
	2		98	14 9	12 5	84	34 4	24 4	71	73	77	115	109
	3		131	14 1	8 6	61	31 2	16 8	54	90	93	68	65
FOMF A	1	ÇC.	ı	14 4	12 3	85	32 3	24 0	74	97	91	88	93
	2		*	-	-	1-	-		-	-	-	-	
	3	"	-	-	-	-	-	-	-	-	-	-	i. —
SOGP - A	1	CC	1	15 0	11 2	75	33 1	21 8	66	83	85	89	87
	2		124	15 5	10 1	65	36 7	19 8	54	54	63	121	103
	3	"	177	19 1	11 1	58	44 7	21 7	48	54	56	107	103
MCP- B	1	CC	2	14 6	11 1	76	33 3	21 7	65	89	82	85	93
c. b	2	"	120	14 6		66	33 0	18 8	57	85	89	77	74
	1 2 3		325	10 2	6 54		22 6	12 8	57	93	93	68	68
DOME D	16	22	2	14.5		7.0	20.1	20.0					
FOMF- B	į,	CC.	2	14 5	11 4	79	32 1	22 2	69	90	84	87	94
	2		135	13 8	10 3	75	31 6	20 1	64	57	67	99	111
	3		190	13 8	6 23	45	31 7	12 1	38	61	66	67	68
SOGP-B	1	CC	2	14 4	11 5	80	63 8	22 4	35	93	93	84	84
a 155	2	"	122	14 2	4 75		32 2		29	78	73	43	46
	3		174	13 8	4 97	36	30 9	9 4 9 7	31	73	81	49	45

^{*} Average Voltage during Discharge = 1 95 volts ** AH+/AHt *** AH+/AH+ dis/AH+

Table IV

SUMMARY DATA-CHARACTERIZATIONS AT MODIFIED CONSTANT POTENTIAL,
FIRST ORDER MASS FLOW, AND SECOND ORDER GAS PRESSURE CONTROLLED CHARGES

Balt No	Char No	Mode	No of Cycles	Amp Charge	ere Hour Dis Eff	Watt Hour Charge Dis* Eff	Percent Charge Acceptance** Pos Neg	AH Recovery (%) Discharge*** Pos Neg
MCP-A	1 2 3	MCP "	97 130	15 0 13 6 14 0	12 5 83 10 1 74 9 4 67	36 / 24 2 66 32 5 20 3 61 32 3 18 3 57	73 75 58 91	102 100 115 74
FOMF-A	1 2 3	FOMF	- -	12 3	10 8 88	29 1 21 0 72 	97 91 	88 93
SOGP-A	1 2 3	SOGP	2 123 176	12 9 7 1 11 7	11 9 92 6 3 89 10 6 91	31 0 23 1 75 15 9 12 3 77 27 0 20 6 76	81 81 84 87 88 89	113 113 105 102 103 101
мср-В	1 2 3	MCP	1 119 324	16 2 10 4 20 0	11 6 71 8 3 80 8 7 44	40 7 22 3 55 24 0 16 2 67 48 0 17 0 35	76 72 95 97 70 58	93 99 84 82 62 75
FOMF-B	1 2 3	FOMF	1 134 189	13 2 9 5 6 8	10 6 80 8 9 94 5 4 79	31 4 20 6 66 22 5 18 5 82 16 2 10 5 6	98 99 89 89 84 84	82 82 106 105 94 94
SOGP-B	1 2	SOGP "	121 173	13 2 11 9 10 4	9 8 74 4 0 30 5 0 48	30 9 19 1 62 27 7 7 7 28 24 6 9 8 40	93 93 79 31 78 76	80 83 42 41 61 63

^{*} Average voltage during discharge | 1 95 volts

Likewise, efficiencies in terms of ampere-hours and watt-hours obtained in discharge relative to the previous charge were about the same for the MCP mode and the two gas-controlled modes. However, each comparison has its own qualification, so that the overall picture is not clear-cut. For example, with respect to the long life of battery MCP-B, this particular battery and charging mode also entailed more undesirable gas evolution than any of the other batteries or methods.

It was also observed that prolonged overchanges at an end of charge rate less than that normally employed result in a capacity recovery for the immediately following discharge. A return to the originally selected end-of-charge current results in an immediate drop in discharge capacity to that observed before the end of discharge current perturbation. This observation shows that the active material is not sloughed off the battery plates or permanently buried under passivated electrode material. It may be due to some other change in the physical structure of the battery plates.

Charging Efficiencies

Columns 7 and 10 of Tables III and IV show that the ampere-hour and watt-hour efficiencies are more determined by the charging mode than by the cycle number in the life of a battery, except insofar as the cycle number reflects a relatively abrupt deterioration of the battery. Thus, the efficiencies in the MCP method decrease slightly during cycle life of batteries MCP-A and MCP-B. This corresponds to a larger fraction of the current going into gas evolution which can only be compensated by an increase in the end-of-charge current. Conversely, in gas control, efficiencies should have been independent of the tendency to-wards gas evolution because a fixed gassing rate controls the charge current. This does not allow undue gassing. Instead, less useful charging takes place. This could be the source of undercharging in the series FOMF-B and SOGP-B.

The most obvious weakness in all batteries was the decrease of hydrogen overvoltage at the negative plates which caused the increased gassing and greater inefficiencies. This is reflected in all three different charging modes. However, inspection of Tables III and IV will show that in terms of ampere-hour efficiency all batteries except FOMF-A and SOGP-B maintained reasonable efficiencies throughout cycle life. FOMF-A failed because of instrumental breakdown, so that SOGP-B was the only poorly performing battery of the six batteries tested.

Speed of Charge

Gas-controlled charges, as they have been applied for this program, have focused on charging efficiency.

^{**} AH+/AHt

^{***} AH_dis/AH-

Table V

Cycle No.	MCP-A Discn. AH	8	FOMF-A Disch.		SOSP-	A. %	MCP-B Disch AH	96	FOMF- Disch AH	-В '3	SOGP-3 Disch
1 2 3 4 5 6 7 8 9	12 5 10 8 11 4 10 9 10 7 10 6 10 7 10 9 10 9		10 8 9 9 12 1 12 4 11 0 11 0 10 7 10 1 11.1		10 8 10 1 10 1 11 9 11 0 11.0 4 9 6 9 2 8 6		11 6 11 4 11 4 11 6 11 4 11 0 10 4 11 1		7 6 7 5 4 7 7 0 7 0 7 1 6 7 6 3 6 0 5 8		9 0 8 3 7 8.1 7 7 6 7 4 7 3 7 4
(Avg 1-10)	(11 0	100)	(11 0	100)	(10 3	100)	(1. 2	100)	(6 6	100)	(8 100)
20 30 40 50 60 70 80 90 100 111	10 4 10 6 10 7 10 9 10 1 9 8 9 8 10 2 11.8	95 96 97 99 92 89 89 93	11 1 10 9 10 7 11 0 10 7 10 0 9 8 9 5 10 2 9 7 9 5	101 99 97 100 97 91 89 86 93 88	11 2 10 3 8 8 7 6 8 6 8 5 8 6 7 0 7 6 7.0	109 100 85 74 83 83 83 68 74 68	11 7 9 6 11 2 8 6 8 9 7 8 8 9 10 8 9 1	104 86 100 77 79 70 79 96 81 39	4 8 8 1 8 0 9 4 9 3 9 2 7 3 9 1	73 73 123 121 142 141 139 1.1 138	6 86 6 87 10 64 3 2 41
120	9 6	87	Failed		6 2	60	9 7	87	9 3	141	4 4 50
129 130 140 150 160 170	5.7	52			6 6 6 1 5.8 5 5 5 7	64 59 56 53 55	9 2 6 7 9 3 9 3 8 7	82 60 83 83 78	8 6 8 9 7 7 6 1 5 2	130 135 1.7 92 79	3 4 3 0 03 5 6 113 5 3 68 4 7 60 4 4 49
175 180					5.5	53	8 8	79	4 7	71	
188 190 200 210 220 230 240 250 260 270 280 290 300 310 320							8 9 8 2 9 7 • 5 5 7 7 6 6 8 8 8 7 6 8 8 8 9 8 6 • 5	79 70 73 71 68 76 65 63	4.8	73	

Where charge acceptance is high, such as at the beginning of charge of a previously completely discharged battery, rate of charging and efficiency of charging are high. As the battery becomes partially charged, its charge acceptance decreases but its charging efficiency remains high by the gas-control methods. This results in a prolonged time of charge at high efficiency because of the diminishingly small amounts of uncharged active material. These data show that, near the beginning of charge, the SOGP method is fastest in terms of charge returned to the battery However, as the charge continues, differences between the SOGP, FOMF, and MCP charges, in terms of capacity restored, become less apparent. By any of the three charging methods, and particularly the gas-controlled methods adjusted for very efficient charging, 100 percent charges of previously discharged capacity are difficult and become more difficult if high charging efficiency is required.

Charge Acceptance and Ampere-Hour Recovery on Discharge

These parameters, given in the last four columns of Tables III and IV, reflect the efficiency of charging individual electrodes and their ability to deliver that charge on the subsequent discharge. Because of the larger amount of gassing in the MCP mode, the charge acceptance in this mode should have been somewhat lower than for the other methods. This was observed in characterization MCP-B-3. Recovery was almost uniformly good except at the end of cycle life in the third characterization and for the second characterization of battery SOGP-B.

MCP Charging

Both batteries MCP-A and MCP-B deteriorated because of usual causes of failure, namely the corrosion of the positive grid, shedding, and bridging of the positive plate materials. In addition, lowering of the hydrogen overpotential at the negative electrode

in batteries with grids containing antimony causes earlier hydrogen gassing, which in turn is reflected in lower charge acceptance. Worsening performance may be due to gassing from the negative plate; the actual battery failure can be attributed to the positive plate. For example, in battery MCP-A, this is consistent with the concurrent decrease of the negative plate potential to lower values during the course of cycle life; viz., 1 41 to 1 22 to 1 14 V vs Hg/HgaSO4 in the three characterizations. Thus, the stoichiometric evolution of hydrogen and oxygen was maintained although the positive material deteriorated to a greater extent than the negative. There was appreciable charge acceptance and good charge retention, limiting gas evolution, in the second characterization, but poor performance in the final characterization.

Battery MCP-B similarly showed a decrease in hydrogen overpotential which required frequent replenishment of water lost by excessive gassing. However, given that handicap, the longest cycle life of all batteries tested was achieved by battery MCP-B.

This also required having the charging cycle terminated at relatively high currents in the later stages of cycling (1.20 to 1.60 amp, equivalent to the C/10 to C/7.5 rates).

GAS-CONTROLLED CHARGE - FOMF AND SOGP

Since these are gas-flow-controlled charges, gassing at lowered negative potential due to antimonial contamination resulted in a concomitant reduction in the charging current. The net effect was a virtually unchanged charging efficiency but longer time on charge as a given battery containing Pb/Sb grids ages. This effect was noted particularly on life-test. Batteries cycled by either the FOMF or the SOGP methods required progressively longer charging times as they aged. For this reason, the end of charge current cutoff was raised resulting in a shorter time on charge.

Previous work had shown the advantages of gascontrolled methods. FOMF and SOGP were selected for further study with respect to life-testing as the most promising methods. However, the life-test data of this study are not sufficiently conclusive to confirm this expectation. Only in the case of SOGP-A was there found a superior charging performance for gas control. As pointed out previously, the longest cycle life was obtained with MCP-B. The advantage of gas control lies in limiting the loss of water from the electrolyte, but this is achieved at the cost of undercharging. An occasional deliberate overcharge of the battery shows that the capacity is still available, but it is not completely used in either of the two gascontrol modes.

FAILURE ANALYSIS

All batteries were subjected to failure analysis. There was less positive grid corrosion and shedding of active material in Type A batteries than in Type B

batteries. SOGP-B failed because of bridging of plates by shedded positive material. Corrosion of positive grids in Type B batteries followed the order MCP > SOGP > FOMF, from worst to least corrosion.

BATTERIES WITH TYPE A VS TYPE B DESIGN

The two types of batteries are mentioned above. Their designs were different. Yet, they are very similar in terms of weight and geometry; i.e., weight and volume of components, and most importantly, as far as the volume, thickness, and surface area of the plates are concerned.

One major difference was the tight wrapping of Type B cell packs in a woven polymeric cloth. This had the advantage of preventing the loosening and shedding of active material, but had also the disadvantage of a more compressed construction causing whatever dendrite or spalling takes place to bridge the plates for a short. In fact, that was observed in at least one case.

Another difference is the composition of the antimonial grid alloy The Type A battery had only about 4 percent Sb in the grid compared to more than 6 percent for the Type B battery This may not be directly related to cycle life or battery performance since antimony caused gassing in both cases. However, inspection of the batteries showed that the positive grids of Type A were less corroded and this may be related to the lower Sb content of Type A grids.

Undercharging occurred in gas-controlled charging modes. It was a more serious problem for Type B than for Type A batteries. The cause of this in terms of battery construction is not known, but the effect was pronounced as shown in Table V.

COMBINATION OF CHARGING MODE WITH BATTERY TYPE

Inspection of Tables III and IV does not show any outstanding difference in the cycle life and performance of the various possible combinations of charging modes and batteries.

Concerning gas-controlled charging, the severe undercharging of batteries FOMF-B and SOGP-B has been mentioned. It was not as much observed with Type A batteries.

Longer life time of MCP-B was obtained. However, this battery exhibited more gassing and experienced more positive grid corrosion than in the corresponding gas control methods. Conversely, MCP-A had a slightly shorter cycle life than SOGP-A and presumably that of FOMF-A, had it not failed for instrumental reasons.

It appears, therefore, that the results of this study do not discriminate strongly between the different charging modes for the kind of batteries and heavyduty cycling regimes used in the present work. Gas control requires more elaborate electronic circuitry Maintenance of precise and sensitive mechanical conditions of gas flow is also necessary. The former presents no problems because there is a very substantial electronic technology, either with operational amplifiers or with miniaturized circuits, which can handle the electrical part of the charging controls. The mechanical control of the gas flow may, however, be more difficult to control in a totally automated system over the lifetime of a battery that is intended to last for many years. Conversely, the MCP method is much simpler in principle; but as detailed above, this method too has its drawbacks.

APPLICATION TO ELECTRIC VEHICLES

Although the batteries used for this program were not traction batteries, the data suggest the applicability of gas-flow-controlled charging for electric vehicles.

Presently, available data relating to the range of road-type electric vehicles power by conventional lead-acid batteries show the need for rapid charging stations particularly on interstate highways and other roads used for relatively long distance travel. There is also an obvious need for overnight charging stations perhaps powered by single-phase 230-volt service normally available in private residences. The data collected during the characterizations of the batteries used for this program illustrate the possibility for quick, efficient charging by the gas-controlled methods and the MCP method where the initial charging currents are at the 2C rates. The same data illustrate that overnight charging would include relatively long time periods at reduced current; a condition found to be desirable in restoring capacity to batteries previously subjected to a series of partial charges.

A convenient method for evaluating a charging system in terms of rapid charging is to measure the percentage of capacity restored in a short time on charge: for example, one-half hour. For this program, the characterization data were compared with the discharge capacity for the discharge immediately preceding the characterization. For new batteries charged by the three methods used for this program, the percentage of previously discharged capacity restored in one-half hour was as follows:

FOMF	63%
MCP	69%
SOGP	90%

In general, as the batteries aged, they recharged as a somewhat diminished but still appreciable rate. This occurred because of reduced charge acceptance; progressively more charging current resulted in gassing.

The above percentage recovery data were obtained on batteries subjected to complete discharges. Other parameters such as a realistic duty cycle, operating temperature, and ultimate battery life need

to be considered before definite conclusions could be reached concerning the effectiveness of a given charging method for quick charging of electric vehicles.

CONCLUSIONS

- 1. The three charging modes, although very different in principle, gave equivalent performances in combination with the lead-acid batteries having antimonial grids.
- 2. The MCP method is preferred for simplicity of equipment and operation. However, more corrosion, more gassing, and the consequently more frequent maintenance were observed and required.
- 3. The gas-controlled methods are relatively maintenance-free. Little water was lost from the batteries by electrolysis, but undercharging was a problem.
- 4. In all cases, more sensitive and uniform control was achieved with gas-controlled charging throughout the cycle life of both types of batteries.
- 5. Charging speed is dependent upon the charging mode mainly in the beginning of charge. As charging progresses, differences in charging rates for the different charging modes become less pronounced.
- 6. Where obvious failure occurred, it was caused by the bridging of positive active material at the edges and at the bottom of the cell pack. Some positive grid corrosion was also noted.
- 7 For electric vehicle application, it appears that a gas-controlled charging system--for example, one depending on differential pressure as the controlling parameter--might be best for quick partial onthe-road charge. This could be combined with a slow, complete, off-the-road overnight charge at reduced current.

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