Regulatory Volume Decrease and Intracellular Ca\textsuperscript{2+} in Murine Neuroblastoma Cells Studied with Fluorescent Probes

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Abstract The possible role of Ca\textsuperscript{2+} as a second messenger mediating regulatory volume decrease (RVD) in osmotically swollen cells was investigated in murine neural cell lines (N1E-115 and NG108-15) by means of novel microspectrofluorimetric techniques that allow simultaneous measurement of changes in cell water volume and [Ca\textsuperscript{2+}], in single cells loaded with fura-2. [Ca\textsuperscript{2+}], was measured ratiometrically, whereas the volume change was determined at the intracellular isosbestic wavelength (358 nm). Independent volume measurements were done using calcein, a fluorescent probe insensitive to intracellular ions. When challenged with 10% hyposmotic solutions, the cells expanded osmotically and then underwent RVD. Concomitant with the volume response, there was a transient increase in [Ca\textsuperscript{2+}], whose onset preceded RVD. For hyposmotic solutions (up to 20%), [Ca\textsuperscript{2+}], increased steeply with the reciprocal of the external osmotic pressure and with the cell volume. Chelation of external and internal Ca\textsuperscript{2+}, with EGTA and 1,2-bis-(o-aminophenoxy) ethane-N,N,N',N''-tetraacetic acid (BAPTA), respectively, attenuated but did not prevent RVD. This Ca\textsuperscript{2+}-independent RVD proceeded even when there was a concomitant decrease in [Ca\textsuperscript{2+}], below resting levels. Similar results were obtained in cells loaded with calcein. For cells not treated with BAPTA, restoration of external Ca\textsuperscript{2+} during the relaxation of RVD elicited by Ca\textsuperscript{2+}-free hyposmotic solutions produced an increase in [Ca\textsuperscript{2+}], without affecting the rate or extent of the responses. RVD and the increase in [Ca\textsuperscript{2+}], were blocked or attenuated upon the second of two 40% hyposmotic challenges applied at an interval of 30–60 min. The inactivation persisted in Ca\textsuperscript{2+}-free solutions. Hence, our simultaneous measurements of intracellular Ca\textsuperscript{2+} and volume in single neuroblastoma cells directly demonstrate that an increase in intracellular Ca\textsuperscript{2+} is not necessary for triggering RVD or its inactivation. The attenuation of RVD after Ca\textsuperscript{2+} chelation could occur through secondary effects or could indicate that Ca\textsuperscript{2+} is required for optimal RVD responses.

Key words: calcium • volume regulation • neuroblastoma • regulatory volume decrease

Introduction Neurons, like all animal cells, possess mechanisms for maintaining constant volume in isosmotic media in the face of the Donnan effect or changes in intracellular solute content, such as those produced by neurotransmitters, hormones, repetitive nerve impulses, or upon sodium pump inhibition (Serve et al., 1988; Alvarez-Leefmans et al., 1992). In addition, neurons, like most cells, regulate their volume when exposed to anisosmotic media (Hoffmann and Simonsen, 1989). Upon exposure to hyposmotnic media, cells initially swell, and then return to their initial volume, a phenomenon termed regulatory volume decrease (RVD).1 This regulatory response is accomplished through the loss of intracellular solutes along with osmotically obligated water. Three components are involved in this negative feedback loop: (a) the volume sensor, which detects a change in cell volume and transduces this change into an intracellular signal; (b) the effector mechanism that causes the loss of osmolutes and water, thereby correcting the aberrant volume; and (c) the signal or second messenger that couples the sensor to the effector. Chief among the candidates for second messenger is intracellular Ca\textsuperscript{2+} (for review see McCarty and O’Neil, 1992). However, the evidence supporting a signaling role for Ca\textsuperscript{2+} is fragmentary and controversial (Grinstein and Smith, 1990; Foskett, 1994). For instance, RVD measured in cell populations of neurons or astrocytes has been claimed to be unaffected by external Ca\textsuperscript{2+} removal, although internal Ca\textsuperscript{2+} was neither chelated nor measured (Pasantes-Morales et al., 1993, 1994). In neuroblastoma cells, RVD occurred in the absence of external Ca\textsuperscript{2+} in cells loaded with the Ca\textsuperscript{2+} chelator 1,2-bis-(o-aminophenoxy) ethane-N,N,N',N''-tetraacetic acid (BAPTA; Lippmann et al., 1995). However, intracellular Ca\textsuperscript{2+} was not measured to verify the effectiveness of Ca\textsuperscript{2+} buffering. A role for Ca\textsuperscript{2+} as mediator of RVD has been

1 Abbreviations used in this paper: AM, acetoxymethyl ester; BAPTA, 1,2-bis-(o-aminophenoxy) ethane-N,N,N',N''-tetraacetic acid; CWV, cell water volume; RVD, regulatory volume decrease; SES, standard external solution.
claimed for various cell types (McCarty and O’Neil, 1992) including astrocytes (O’Connor and Kimelberg, 1993) and cortical neurons (Churchwell et al., 1996). More recently, it was concluded that Ca\(^{2+}\) signaling by Ca\(^{2+}\) release or Ca\(^{2+}\) entry appears to play no role in the activation mechanism for the RVD responses in Ehrlich cells (Jørgensen et al., 1997). However, changes in intracellular free Ca\(^{2+}\) concentration, [Ca\(^{2+}\)], were measured in cell populations or on single cells, in which changes in volume were not measured simultaneously. Clearly, part of the above controversy is due to the fact that changes in [Ca\(^{2+}\)], and volume have not been measured simultaneously in single cells and a causal relationship between these two variables has not been definitively established in the above studies.

In the present study, we unify the above conflicting claims in the literature by directly demonstrating that RVD can proceed independently of a rise in intracellular Ca\(^{2+}\), but that there is a component of the response that is sensitive to external and internal Ca\(^{2+}\) chelation. We used novel optical techniques to simultaneously measure changes in [Ca\(^{2+}\)], and cell volume in response to hyposmotic challenges in two neuronal cell lines (N1E-115 and NG108-15). Changes in cell water volume (CWV) were estimated from the change in intracellular concentration of trapped fluorescent dyes (Alvarez-Leefmans et al., 1995). By using the Ca\(^{2+}\)-sensitive dye fura-2 and recording at the Ca\(^{2+}\)-sensitive (380 nm) and Ca-sensitive (isosbestic, 358 nm) wavelengths, we monitored simultaneous changes in CWV and [Ca\(^{2+}\)], in single cells (Muallem et al., 1992). CWV changes were also measured using calcine as the fluorescent probe (Crowe et al., 1995), thus providing independent measurements of CWV with a dye having different chemical properties than fura-2.

We find that upon hyposmotic swelling there is a rise in [Ca\(^{2+}\)], whose onset precedes RVD. However, chelation of external and internal Ca\(^{2+}\) only attenuates, but does not prevent, RVD. Moreover, this Ca\(^{2+}\)-independent RVD proceeds even when [Ca\(^{2+}\)] decreases below resting level. We also find that upon multiple hyposmotic challenges the RVD response is lost. We call this effect inactivation. This phenomenon is also independent of Ca\(^{2+}\). Two models are consistent with our observations. In the first model, a single sequential process underlies RVD. In this model, a rise in [Ca\(^{2+}\)] cannot be the triggering signal for RVD. The attenuation of RVD after Ca\(^{2+}\) chelation could occur through secondary effects or could indicate that Ca\(^{2+}\) is required for optimal responses. In the second model, RVD could be the consequence of the operation of two parallel processes, one of which is Ca\(^{2+}\) independent. Some of these results have been reported in preliminary form (Altamirano et al., 1996).

### Materials and Methods

#### Cell Culture

Two murine neural cell lines maintained in culture, N1E-115 neuroblastoma and neuroblastoma x glioma NG108-15, were used in the present experiments. N1E-115 is a clone of cells derived from mouse neuroblastoma C-1300 (Amano et al., 1972), and expresses characteristics of neuronal cells. The neuroblastoma x glioma hybrid cells NG108-15 are hybrid cells obtained from a rat glioma cell line and a mouse neuroblastoma line, and express characteristics of both glial and neuronal cells (Hamprecht, 1977). Cells were grown at 37°C on culture dishes containing 90% Dulbecco’s modified Eagle’s medium (DMEM) supplemented with 10% FCS, 1% hypoxanthine-aminopterin-thymidine (HAT) and 1% L-glutamine, in a 5% CO\(_2\)/95% air atmosphere. Cells from passages 9–25 were plated on 25-mm-diameter glass coverslips (Bellco Glass, Inc., Vineland, NJ), previously treated with poly-lysine. Differentiation was induced 24 h after plating by supplying the cells with a low serum growth medium composed of 98% DMEM, 1% FCS, 1% HAT, 1% L-glutamine, 1 mM theophylline, and 10 μM prostaglandin E\(_1\) (Kasai and Neher, 1992). Cells were used from 1 to 14 d after the differentiation treatment.

#### Saline Solutions

The standard external solution (SES) contained (mM): 130 NaCl, 5.5 KCl, 2.5 CaCl\(_2\), 1.25 MgCl\(_2\), 20 HEPES, 10 dextrose, 13 sucrose. The pH was adjusted to 7.3 with NaOH and the osmolality was 312 ± 3 mosmol/kg water. The control isosmotic solution was prepared by substituting 65 mM NaCl for sucrose to match the osmolality of the culture medium (312 mosmol/kg water). Anisosmotic solutions were prepared by sucrose addition or removal to get the desired osmolality, thus maintaining constant the ionic strength and at the same value as the isosmotic control. Anisosmotic solutions were expressed as the decrement (or increment) percentage with respect to the control isosmotic solutions. Thus, a solution referred to as a “44% hyposmotic” means that its measured osmolality was 56% of the control isosmotic solution. The zero Ca\(^{2+}\) solutions were made by substituting CaCl\(_2\) for MgCl\(_2\) and adding 0.5 mM EGTA. The osmolality of all solutions was measured with a vapor pressure osmometer (5100 B; Wescor Inc., Logan, UT).

#### Dye Loading and Bath Chamber

Each coverslip with the cells attached was mounted in a Leiden chamber (Medical Systems Corp., Greenval, NY) filled with SES and placed on the stage of an epifluorescence inverted microscope. Cells were loaded with the fluorophores fura-2 or calcine as described previously (Alvarez-Leefmans et al., 1995). In brief, they were incubated at room temperature (22°C) in SES containing either 5 μM fura-2 acetoxymethyl ester (AM), or 2 μM calcine/AM. To prepare the fura-2-loading solution, we used a stock containing 50 μg fura 2/AM dissolved in 4 μl DMSO plus 4 μl pluronic (10% wt/wt in DMSO). The final concentration of fura-2/AM for this stock solution was 6.23 mM. The calcine loading solution was prepared from a stock containing 50 μg calcine/AM dissolved in 4 μl DMSO plus 4 μl pluronic (10% wt/wt in DMSO). The final calcine/AM concentration of this stock solution was 6.28 mM. Dye loading was monitored fluorometrically by sampling a single cell every 30 or 60 s until reaching the desired level of fluorescence. For calcine, the level was =14.3× the initial fluorescence (i.e., the cell fluorescence without dye), and for fura-2 the level was =8, at 358 nm. The loading time was 37 ± 3
min for calcein \((n = 22)\) and \(43 \pm 3\) min for fura-2 \((n = 26)\). The loading solution was then washed out with SES for at least 1 h before starting the experimental measurements. All solutions, except those used for loading (see below), were perfused into the chamber at a rate of 3 ml/min. The chamber fluid was exchanged with a time constant of \(3.6 \pm 0.3\) s.

**BAPTA Loading**

To buffer intracellular \(\text{Ca}^{2+}\) near resting levels, in some experiments we used the \(\text{Ca}^{2+}\) chelator BAPTA. In this series of experiments, after the cells were loaded with either calcein or fura-2 as described above, they were briefly washed with SES and loaded with BAPTA. This was accomplished by incubating the cells at room temperature for 2 h in SES containing 100 \(\mu\)M BAPTA-AM. The BAPTA loading solution was prepared from a stock containing 3.82 mg of BAPTA/AM dissolved in 1 ml DMSO. After BAPTA loading, cells were superfused with SES for 1 h before starting the experimental measurements. The chelating action of intracellular BAPTA was verified by ratiometric measurements with fura-2.

**Measurement of Cell Volume with Fluorescent Dyes**

**Basic principle.** Changes in CWV can be assessed by measuring changes in concentration of impermeant substances (volume markers) introduced into cells (Alvarez-Leefmans et al., 1995). Intracellularly trapped fluorescent dyes such as calcein (Crowe et al., 1995) and fura-2 (Muallem et al., 1992) have been successfully used as volume indicators. These optical techniques provide superior time resolution and sensitivity relative to other techniques and, when using ratiometric dyes, they allow parallel measurements of changes in CWV and intracellular ion concentrations in single cells. Changes in CWV are estimated from changes in fluorescence intensity of the dye resulting in turn from changes in its intracellular concentration. By using the \(\text{Ca}^{2+}\)-sensitive dye fura-2 and recording at the \(\text{Ca}^{2+}\)-sensitive \((380\) nm\) and -insensitive (isosbestic) wavelengths, we recorded simultaneously changes in CWV and \([\text{Ca}^{2+}]\), in single cells. CWV changes were inferred from changes in the concentration of fura-2, recorded at the isosbestic wavelength (see below). The simultaneous changes in \([\text{Ca}^{2+}]\), were monitored by taking the ratio 358/380. Independent and parallel measurements of CWV were provided by using calcein, a dye that is more intensely fluorescent than fura-2 and insensitive to changes in the concentration of native cellular ions within physiological ranges. The validation, pitfalls, and limitations of these techniques have been discussed in detail in previous publications (Alvarez-Leefmans et al., 1995; Crowe et al., 1995, 1996).

**Fluorescence measurements.** Total fluorescence from a small region of fluorophore-loaded single cells was measured using a customized microspectrophotometry system that has been described in detail elsewhere (Alvarez-Leefmans et al., 1995). The system included an inverted, epifluorescence microscope (Diaphot-TMD; Nikon, Tokyo, Japan) equipped with a fluor oil-immersion objective lens \((0.1–10\) \(\mu\)m). The excitation light coming from a 150 W xenon arc lamp passed through a water filter, and then was divided by a beam splitter. Each beam passed through a computer controlled high speed shutter (Uniblitz; Vincent Associates, Rochester, NY). Attached to each shutter was a filter wheel holding a selection of customized excitation filters (Omega Optical, Brattleboro, VT). For fura-2 measurements, beam 1 passed through a filter centered at 380 ± 6 nm. For exciting fura-2 at its isosbestic wavelength, beam 2 passed through a filter centered at 362 or 358 ± 5 nm. The 358-nm wavelength was preferred over the 362-nm because it was closer to the intracellular isosbestic wavelength measured in vivo with a monochromator and the same objective lens used in the present work (Crowe et al., 1996). In some experiments using the monochromator, measurements were done at wavelengths of 350 and 358 nm \((slit = 8\) nm\) to optimize the sampling rate.

Optical systems may distort spectral responses of fluorescent dyes through various components in the light path; e.g., the objective lens having different spectral light transmission properties or the bandwidth of the excitation light. Fig. 1 shows a bath calibration of fura-2 performed in the recording chamber with the optical system used in the present experiments. This calibration shows that at 358 nm, the fura-2 signal was insensitive to changes in \([\text{Ca}^{2+}]\) in the range between 0 and 39.80 \(\mu\)M. As expected, the dye showed adequate sensitivity for \(\text{Ca}^{2+}\) at the 380-nm wavelength. The latter wavelength was preferred over 340 nm because the light transmission properties of the objective used were better at 380 nm. Light transmission of this objective is 74% at 380 nm, 68% at 360 nm, and 55% at 340 nm.

For CWV measurements using calcein, beam 1 passed through a filter centered at 495 ± 10 nm to excite this fluorophore at its peak excitation wavelength. When dual excitation wavelength measurements were done (fura-2), the filtered light beams were recombined by means of a bifurcated, fused silica-fiber bundle. The combined light beam was then collimated and coupled to the microscope epi-illuminator. To minimize photobleaching and photodynamic damage, the intensity of the excitation light beam was attenuated with a 10% neutral density filter and the shutter was opened only during data sampling (see below). The field diaphragm was closed to its smallest opening to limit the excitation area to a single cell. For fura-2 measurements, a cube holding a dichromatic mirror \((400\) nm\) and a 510 ± 20 nm emission filter was positioned underneath the objective lens in the filter cassette holder of the microscope. For calcein, a dichromatic mirror \((515\) nm\) and a 535 ± 13 emission filter was used instead. The emitted light passed through a one-times relay lens coupled to a turret diaphragm accessory (Microflex PFX; Nikon) containing a series of seven circular pinhole openings of varying diameters \((0.1–10\) mm\).

![Figure 1. Common mode of rejection test. Fura-2 fluorescence recorded in the experimental bath chamber in response to a series of 0–10 mM calcium-EGTA buffer solutions, using a fluor 40×/1.3 oil-immersion objective. F, fluorescence in arbitrary units (a.u.). The free [Ca\(^{2+}\)] (micromolar) is indicated by vertical arrows. Fura-2 was excited at 358 and 380 nm. Emission was measured at 510 nm. Note that at 358 nm, the isosbestic wavelength, the dye does not respond to Ca\(^{2+}\).](jgp.rupress.org)
The emitted light exiting through a selected pinhole (see below) was measured with a photometer system as described previously (Alvarez-Leefmans et al., 1995). Once amplified, the photomultiplier signal was digitized with an analog to digital converter, displayed, and stored on an IBM-compatible PC computer using customized software. The fluorescence data were collected at a frequency of either 0.1 or 0.2 Hz during shutter openings of 130-ms duration. Subsequent data analysis and presentation were performed using commercial software.

As CNV changes, the concentration of osmotically active fluorophore molecules changes in inverse proportion. To observe changes in fluorescence intensity due to changes in intracellular fluorophore concentration, it is necessary to record from a small region of the loaded cell. With epi-illumination, the image of the region of the cell from which light is measured is a volume element determined by the pinhole aperture placed at the image plane and the numerical aperture of the objective. The pinhole aperture used for collecting calcine fluorescence was 0.2 mm in diameter and, with the objective lens used, the combination gave a total recording area of 19.6 µm². The mean area of the cells, at the image plane, was between 1,250 and 1,950 µm². The pinhole used for measuring fura-2-emitted fluorescence was 0.5 mm in diameter, giving a total recording area of 122.7 µm². The larger pinhole is necessary to improve the signal/noise ratio in fura-2 recordings. This is because the fura-2 fluorescence is dimmer than that emitted by calcine, as expected from their corresponding quantum yields and extinction coefficients (Haugland, 1996), and also because of the spectral transmission properties of most objective lenses (Keller, 1995), including the one we used (see above). We have studied the effects of pinhole size and location on the signals recorded at 358 nm in fura-2-loaded cells. It was found that there is a compromise between pinhole size, resolution, and signal-to-noise (S/N) ratio. We have determined that optimal S/N ratio with loss of resolution is obtained with a pinhole located well within the cell boundaries and comprising no more than 10% of the total area of the cell body (Crowe et al., 1996).

Calculation of cell water volume changes. Normalized cell water volume changes (Vt/Vo) were computed from monitored changes in relative fluorescence (F/Fo) resulting from exciting fura-2 at 358 nm or calcine at 495 nm, according to the following equation (Alvarez-Leefmans et al., 1995):

\[
\frac{F_t - F_{bkg}}{1 - F_{bkg}} = \frac{V_t}{V_o},
\]

where Ft is the fluorescence from a pinhole region of the cell equilibrated with an isotonic SES or in a solution isosmotic and isotonic with the SES, having an osmotic pressure πo; Ft is the fluorescence of the same region of the cell in a solution of osmotic pressure πt; Fbkg is the background fluorescence (see below); Vt is the water volume of the cell in a solution having an osmotic pressure πt and Vt is the corresponding volume of the cell in a solution of osmotic pressure πt. In the present study, Fbkg includes the intrinsic cell fluorescence plus the osmotically insensitive component of the fluorescence in dye-loaded cells. Both components of Fbkg are insensitive to changes in external osmolality. Fbkg was estimated from the y-intercept of plots of Ft/Fo vs. πt/πo, as described previously (Alvarez-Leefmans et al., 1995). It is worth noting that Fbkg determined graphically is similar to the fluorescence remaining after controlled plasma membrane permeabilization with α-toxin (Crowe et al., 1995). Moreover, the two approaches have been validated by direct morphometric measurements of cell volume using differential interference contrast (Alvarez-Leefmans et al., 1995) or confocal image reconstruction (unpublished observations).

We characterized the relation between Ft/Fo vs. πt/πo for each individual cell. Steady state changes in Ft/Fo were measured in response to brief (5–10 min) exposures to anisosmotic solutions having nominal osmolalities in the range between ~10 and +10% relative to the isosmotic solution. Within this range of external osmolalities and exposure times, the cells exhibited osmotic behavior and did not regulate their volume. Moreover, the peak response to hypotonic solutions having osmolalities ~40% was also osmometric before the cells started undergoing RVD. All experiments were done at room temperature (22°C).

Drift Correction and Statistical Analysis

When necessary, the drift of the fluorescence signal resulting from dye leakage and photobleaching was corrected by fitting a linear regression to the base line and multiplying the slope of this regression line by the time at which each data point was sampled. This process yields point by point drift values that, in turn, are subtracted from the record yielding the corrected trace.

RVD response variables were measured as defined in Fig. 2. The extent of RVD (Fig. 2 f) was measured 20 min after the onset of the regulatory response and is presented as % RVD:

\[
\% \text{RVD} = \left( \frac{\text{peak } V_t}{V_o} - 1 \right) - \left( \frac{\text{reg } V_t}{V_o} - 1 \right) \times 100
\]

where peak Vt/Vo and reg Vt/Vo are the peak and regulated relative volumes, respectively. Therefore, % RVD denotes the magnitude of the return from the peak swollen volume back toward the isosmotic initial volume, V0, such that 0% RVD indicates no volume regulation and 100% RVD indicates complete volume recovery. Referring to Fig. 2, % RVD is simply \[(c - a)/a \] × 100. All measurements are expressed as mean ± SEM. The one-tailed un-

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**Figure 2.** Schematic drawing illustrating measured variables of osmotic swelling and RVD responses elicited by a 40% hypotonic challenge. Ordinate: relative cell volume (V/V0). Abscissa: time. (a) Maximum (peak) swelling, (b) initial rate of swelling, (c) time to peak swelling, (d) RVD delay, (e) initial rate of RVD, (f) extent of RVD at t = 20 min, and (g) regulated relative cell volume at t = 20 min.
paired t test was used to establish the significance of differences between means at the 95% confidence limit.

Chemicals

All chemicals used to prepare bathing media came from Sigma Chemical Co. (St. Louis, MO). Tissue culture media (Dulbecco’s modified Eagle’s medium), L-glutamine, and hypoxanthine-aminopterin-thymidine came from GIBCO BRL (Gaithersburg, MD), FCS from HyClone (Logan, UT), theophylline and prostaglandin E1 from Sigma Chemical Co., and poly-D-lysine from ICN (Costa Mesa, CA). Fura-2/AM, fura-2-free acid, calcein/AM, BAPTA/AM pluronic, and calcium buffers came from Molecular Probes, Inc. (Eugene, OR).

Results

Regulatory Volume Decrease in Single Cells Loaded with Calcein or Fura-2

Regulatory volume decrease was studied in cultured N1E-115 and NG108-15 cells loaded with either calcein or fura-2. This strategy provided an independent measurement of CWV with dyes having different chemical properties, thus giving more reliability to our observations. In particular, calcein fluorescence is independent of changes in the concentration of native intracellular ions within the physiological range. The parallel use of calcein eliminates possible intracellular Ca2+ buffering effects of fura-2, which may attenuate putative Ca2+-dependent components of the RVD response pattern. On the other hand, the use of fura-2 allowed for the simultaneous measurement of changes in [Ca2+]i and CWV in single cells.

Fig. 3 shows a typical RVD pattern in response to a hyposmotic challenge recorded in a single NG108-15 cell loaded with calcein. The cell was initially superfused with an isosmotic solution to obtain a stable baseline. Upon exposure to a 44% hyposmotic solution, the calcein relative fluorescence (F/F0) decreased due to intracellular dye dilution resulting from net osmotic influx of water. After reaching a minimum, the F/F0 signal recovered close to initial values, reflecting a gradual increase in intracellular dye concentration resulting from regulatory net water efflux. When the hyposmotic solution was replaced with the isosmotic solution, the F/F0 signal increased beyond control levels, reached a peak, and recovered to initial values. This last sequence of fluorescence signals was also expected because, upon returning to the isosmotic solution from a hyposmotic one, the isosmotic solution is hypertonic with respect to the cell that shrinks, and then recovers its initial water volume. Fluorescence signals like these were transformed into relative cell water volume changes (Vt/V0) using Eq. 1. This transformation yielded traces like the one shown in the bottom of Fig. 3. Before testing the 44% hyposmotic solution, the cell was exposed

![Figure 3](https://www.jgp.org/cgi/content/full/55/3/1169/DC1)

Figure 3. RVD monitored in a calcein-loaded NG108-15 cell in response to a 44% hyposmotic challenge. (top) Percent change in relative fluorescence (F/F0). (bottom) Relative cell water volume changes (Vt/V0) computed using Eq. 1. The box indicates the time of application of isosmotic (iso) and hyposmotic (hypo) solutions. The cell swelled to a maximum (83% of its initial volume) and then underwent RVD. Maximal volume recovery (87%) was achieved 22.7 min after RVD onset. As expected, upon returning to the isosmotic solution, the cell shrank and eventually recovered its initial volume. (inset) Vt/V0 in response to calibration test anisosmotic solutions having nominal osmolalities of ±10% with respect to the isosmotic solution (actual osmolalities were −12 and +10%). Steady state Vt/V0 changes in response to the test solutions are plotted against the reciprocal of the relative osmotic pressure of the medium (π0/πt) for the calibration pulses (●). The peak Vt/V0 produced by the 44% hyposmotic solution is also plotted (○). The solid line denotes the theoretical behavior of a perfect osmometer according to Eq. 3. This kind of plot was made for all cells and is shown in subsequent figures.
to calibration test solutions (Fig. 3, inset). These calibration solutions had osmolalities of +10 and −12% with respect to the isosmotic solution. The inset includes a plot that was routinely made for each cell, showing the relationship between steady state values of changes in \( V_t/V_o \) and the reciprocal of the relative osmotic pressure of the medium (\( \pi_o/\pi_t \)). The line denotes the predicted behavior of a perfect osmometer according to the equation:

\[
\frac{V_t}{V_o} = \frac{\pi_o}{\pi_t}
\]

where \( V_o, \pi_o, \) and \( \pi_t \) have been defined. (Fig. 3, ●) Osmotic calibration pulses. Note that the measurements fall close to the theoretical line for a perfect osmometer. (Fig. 3, ○) The peak amplitude of the osmotic response produced by the 44% hyposmotic solution, which also falls close to the theoretical line denoting ideal behavior, before RVD ensues.

Experiments like the one described were performed in seven NG108-15 cells. Upon exposure to a solution ~40% hyposmotic (41.3 ± 1%), the cells swelled to a maximum of 69 ± 5% above their initial volume, which is not significantly different from the value of 71 ± 3% predicted for ideal behavior. The time to reach the maximum swelling was 4.4 ± 0.5 min. After a delay of 47 ± 6 s measured from the peak of the osmotic swelling, RVD ensued at an average initial rate of −3.4 ± 0.6% min\(^{-1}\) with a partial volume recovery at 20 min (percent RVD recovery) of 57 ± 11%. Upon returning to the control isosmotic solution (which would now be hypertonic with respect to the cells), the cells shrank and their volume started to recover. In two cells, complete recovery occurred at 37 and 103 min, respectively, upon returning to the isosmotic solution. The above results are summarized in Table I. Similar results were obtained in three N1E-115 neuroblastoma cells (see Crowe et al., 1995).

The prevailing controversy about the possible role of intracellular Ca\(^{2+}\) as the signal that couples the activation of the volume sensor and the transport mechanisms for RVD prompted us to measure simultaneously \([Ca^{2+}]_i\), and volume changes in fura-2-loaded cells. Fig. 4 shows that a 42% hyposmotic solution produced an increase in \([Ca^{2+}]_i\), accompanied by RVD. When exposed to an ~40% hyposmotic solution (41.4 ± 0.6%), fura-2-loaded NG108-15 cells (\( n = 10 \)) swelled to a maximum of 59 ± 5% above their initial volume. The time to reach the maximum swelling was 5.0 ± 0.4 min. After a delay of 90 ± 23 s, RVD ensued at an average initial rate of −3.0 ± 0.5% min\(^{-1}\) with partial recovery (percent RVD) of 45 ± 8%. The ratio 358/380, which signals changes in \([Ca^{2+}]_i\), started to increase with a latency of 29.5 ± 12 s measured from the onset of the volume change produced by the hyposmotic challenge. The ratio peaked at 7.6 ± 1.4 min after the onset of the volume change produced by the hyposmotic challenge; i.e., ~1 min after the onset of RVD. Similar results were obtained in N1E-115 cells (\( n = 4 \)) following the same protocol. However, in cells loaded with fura-2, the mean values of some variables of RVD responses with respect to those recorded in calcine-loaded cells were slightly attenuated (e.g., maximum swelling and percentage of RVD) or slowed down (e.g., time to peak swelling), but the differences were not statistically significant (Table I).

These results show that fura-2-loaded cells behave like those loaded with calcine in terms of their response to anisomotic solutions and that the change in

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**Table I**

<table>
<thead>
<tr>
<th>Variable</th>
<th>Calcine</th>
<th>Fura-2</th>
<th>( P )</th>
</tr>
</thead>
<tbody>
<tr>
<td>Osmolality (%)</td>
<td>n = 7</td>
<td>n = 10</td>
<td></td>
</tr>
<tr>
<td>Peak swelling (%)</td>
<td>−41.3 ± 1</td>
<td>−41.4 ± 0.6</td>
<td>NS</td>
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<tr>
<td>Observed</td>
<td>69 ± 5</td>
<td>59 ± 5</td>
<td>NS</td>
</tr>
<tr>
<td>Osmometric</td>
<td>71 ± 3</td>
<td>71 ± 2</td>
<td>NS</td>
</tr>
<tr>
<td>Time to peak (min)</td>
<td>4.4 ± 0.5</td>
<td>5.0 ± 0.4</td>
<td>NS</td>
</tr>
<tr>
<td>RVD latency (s)</td>
<td>47 ± 6</td>
<td>90 ± 23</td>
<td>NS</td>
</tr>
<tr>
<td>RVD rate (% min(^{-1}))</td>
<td>−3.4 ± 0.6</td>
<td>−3.0 ± 0.5</td>
<td>NS</td>
</tr>
<tr>
<td>% RVD</td>
<td>57 ± 11 (( n = 6 ))</td>
<td>45 ± 8 (( n = 7 ))</td>
<td>NS</td>
</tr>
</tbody>
</table>

Values are means ± SEM.

---

**Figure 4.** Relative volume changes (\( V_t/V_o \)) and \([Ca^{2+}]_i\), recorded simultaneously in response to a 42% hyposmotic challenge in a fura-2-loaded NG108-15 cell. (top) \( V_t/V_o \) obtained from changes in fluorescence measured at 358-nm excitation wavelength, the isosmotic point for fura-2. (bottom) Changes in \([Ca^{2+}]_i\), measured as the ratio 358/380. Dashed vertical lines are traced at the onset (1) and peak (2) of the volume response, and at the peak of the intracellular Ca\(^{2+}\) signal (3). (inset) Calibration plot obtained as explained in Fig. 3.
cell volume is parallel to a rise in $[\text{Ca}^{2+}]_i$. Moreover, the temporal relationship between changes in $[\text{Ca}^{2+}]_i$ and RVD is appropriate for a putative signaling role of internal Ca$^{2+}$. However, the results do not prove a causal relation between the rise in $[\text{Ca}^{2+}]_i$ and RVD.

**Relationship between External Osmolality, $[\text{Ca}^{2+}]_i$, and Cell Volume**

To test further the Ca$^{2+}$ hypothesis for RVD, we recorded simultaneously changes in volume and $[\text{Ca}^{2+}]_i$ in cells loaded with fura-2 that were exposed to solutions of various osmolalities. Fig. 5 A shows an example in an NG108-15 cell. No measurable changes in the $[\text{Ca}^{2+}]_i$ signal were produced by a 14% hyperosmotic solution that caused an osmometric shrinkage. For hyposmotic solutions, $[\text{Ca}^{2+}]_i$ increased steeply with $p_o/p_t$ (Fig. 5 B, I and II). The rate of RVD increased with $[\text{Ca}^{2+}]_i$ and $V_t/V_o$. The peak of the responses I and II fall close to the value predicted for the behavior of a perfect osmometer (Fig. 5 B, I and II), thus confirming that RVD started after the cell had increased its water volume in an osmometric manner. The above findings show again a close parallel between changes in $[\text{Ca}^{2+}]_i$, cell volume, and rate of RVD. However, they do not prove that the increase in $[\text{Ca}^{2+}]_i$ upon osmotic swelling is a necessary causal signal for RVD and not just an independent parallel process.

**Changes in $[\text{Ca}^{2+}]_i$ and RVD upon Repeated Hyposmotic Challenges**

The close parallel between changes in $[\text{Ca}^{2+}]_i$ and RVD was further shown by exposing the cells to two 40% hyposmotic challenges, each lasting $\sim$40 min, applied...
with an interval of 30–60 min. RVD and the concomitant increase in [Ca\(^{2+}\)], did not occur or were significantly attenuated upon exposure to the second hyposmotic challenge. Fig. 6 illustrates one of these experiments performed in an N1E-115 cell. The first hyposmotic challenge produced the expected transient increase in [Ca\(^{2+}\)], accompanied by RVD. However, the second hyposmotic challenge, applied 55 min after the end of the first one, produced a measurable increase in neither [Ca\(^{2+}\)], nor RVD. In some experiments, the second hyposmotic challenge produced an increase in [Ca\(^{2+}\)], albeit smaller than the one evoked by the first hyposmotic challenge, without a concomitant RVD response. The disappearance of the RVD response upon the second hyposmotic shock was also observed in calcein-loaded cells in both cell lines (Fig. 7).

To test for the external Ca\(^{2+}\) dependence of RVD inactivation, calcein-loaded cells were exposed to Ca\(^{2+}\)-free isosmotic and hyposmotic solutions containing 0.5 mM EGTA. The cells were first superfused with the Ca\(^{2+}\)-free isosmotic solution for at least 10 min, and then the Ca\(^{2+}\)-free 40% hyposmotic solution was applied twice with an interval of ~50 min. The results show that both RVD and its inactivation upon the second hyposmotic challenge proceeded in the virtual absence of external Ca\(^{2+}\) (Fig. 7).

We conclude that RVD inactivation is independent of (a) the type of dye with which the cells are loaded, (b) the cell type (NG108-15 or N1E-115), and (c) the presence of external Ca\(^{2+}\).

The fact that RVD disappearance coincides with suppression or attenuation of the concomitant change in [Ca\(^{2+}\)], confirms the parallel between a rise in [Ca\(^{2+}\)], and RVD.

**Effect of Extracellular and Intracellular Ca\(^{2+}\) Chelation on RVD and [Ca\(^{2+}\)],**

The experiments so far described demonstrate a tight correlation between changes in [Ca\(^{2+}\)], and RVD. Demonstration of this correlation is necessary but not sufficient to suggest a causal role for Ca\(^{2+}\) as a second messenger mediating RVD. A crucial test for the Ca\(^{2+}\) hypothesis is to show directly that RVD is abolished when the rise in intracellular Ca\(^{2+}\) upon osmotic swelling is inhibited. Elevations of [Ca\(^{2+}\)], produced by hyposmotic solutions could result either from Ca\(^{2+}\) influx across the plasma membrane or Ca\(^{2+}\) release from internal stores (McCarty and O’Neil, 1992; Wu et al., 1997). Thus, we studied the effect of extracellular and intracellular Ca\(^{2+}\) chelation on RVD. To remove the external source of Ca\(^{2+}\), these experiments were done with Ca\(^{2+}\)-free EGTA isosmotic and hyposmotic solutions. Intracellular Ca\(^{2+}\) was chelated with BAPTA. For this purpose, cells were incubated with 100 \(\mu\)M BAPTA/AM for 2 h, loaded with calcein, and then su-

![Figure 6. Inactivation of RVD response upon repeated hyposmotic challenges. Changes in relative cell volume (\(V/V_0\)) and [Ca\(^{2+}\)], in response to two identical hyposmotic challenges in a fura-2-loaded N1E-115 cell. The cell was exposed to two 40% hyposmotic challenges, each lasting ~40 min applied with an interval of 55 min. (top) \(V/V_0\) (bottom) Changes in [Ca\(^{2+}\)]. (inset) Calibration plot obtained as explained in Fig. 3. (○) Peak amplitude of the first pulse, and (△) peak amplitude of the second pulse. Boxes indicate the time of application of isosmotic (iso) and hyposmotic (hypo) solutions.](image-url)
perfused with the Ca\(^{2+}\)-free EGTA isosmotic solution for at least 10 min before the Ca\(^{2+}\)-free EGTA hyposmotic solution. The use of calcein was important in these experiments because this dye does not affect intracellular Ca\(^{2+}\) and does not have the side effects of fura-2. We compared a control population of cells with those treated with BAPTA and Ca\(^{2+}\)-free EGTA solutions. Such Ca\(^{2+}\) chelation did not abolish RVD; however, there was a significant (P < 0.01) decrease in the rate (53%) and extent (56%) of RVD responses (Table II).

The persistent RVD could be due to incomplete intracellular Ca\(^{2+}\) chelation. The ability of BAPTA to chelate intracellular Ca\(^{2+}\), like that of other Ca\(^{2+}\) chelators, depends on the [Ca\(^{2+}\)]\(_i\), the ionic strength and composition of the cytosol, and the [BAPTA]. Unless one knows these factors, it is not possible to be sure that BAPTA is adequately chelating internal Ca\(^{2+}\). In the absence of such knowledge, it is essential to verify that BAPTA is adequately chelating intracellular Ca\(^{2+}\). To verify that BAPTA was actually damping any hypsomotically induced increases in [Ca\(^{2+}\)]\(_i\), another series of experiments was done with fura-2-loaded cells. Under these conditions, the hypsomotopic challenge resulted in a decrease in [Ca\(^{2+}\)]\(_i\) below resting levels, but the RVD response persisted (Fig. 8).

### Table II

<table>
<thead>
<tr>
<th>Variable</th>
<th>Control Ca(^{2+})-free</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>Osmolality (%)</td>
<td>41.3 ± 1.0</td>
<td>41.3 ± 0.8</td>
</tr>
<tr>
<td>Peak swelling (%)</td>
<td>69 ± 5</td>
<td>68 ± 4</td>
</tr>
<tr>
<td>Osmometric</td>
<td>71 ± 3</td>
<td>71 ± 2</td>
</tr>
<tr>
<td>Time to peak (min)</td>
<td>4.4 ± 0.5</td>
<td>5.6 ± 0.9</td>
</tr>
<tr>
<td>RVD latency (s)</td>
<td>47 ± 6</td>
<td>73 ± 31</td>
</tr>
<tr>
<td>RVD rate (% min(^{-1}))</td>
<td>-3.4 ± 0.6</td>
<td>-1.6 ± 0.3</td>
</tr>
<tr>
<td>% RVD</td>
<td>57 ± 11 (n = 6)</td>
<td>25 ± 4 (n = 6)</td>
</tr>
</tbody>
</table>

Values are means ± SEM.

---

**Figure 7.** Inactivation of RVD response upon repeated hypsomotic challenges in the virtual absence of external Ca\(^{2+}\). Changes in relative cell volume (V/\(V_0\)) in response to two hypsomotic challenges of identical osmolality, in a single NG108-15 cell loaded with calcein. Except for the calibration test solutions, the isosmotic and hypsomotic solutions were Ca\(^{2+}\)-free and contained 0.5 mM EGTA. After testing the anisosmotic calibration pulses, the cell was superfused with a Ca\(^{2+}\)-free isosmotic solution for 10 min before the first Ca\(^{2+}\)-free 40% hypsomotic challenge. The second Ca\(^{2+}\)-free hypsomotic challenge was applied 44 min after the end of the first one. During the interval between hypsomotic challenges, the cell was superfused with Ca\(^{2+}\)-free isosmotic solution. The inset shows a calibration plot obtained as explained in Fig. 3. (○) Peak amplitude of the first pulse, and (△) peak amplitude of the second pulse. Boxes indicate the time of application of isosmotic (iso) and hypsomotic (hypo) solutions.

**Figure 8.** Effect of extracellular and intracellular Ca\(^{2+}\) chelation on RVD and [Ca\(^{2+}\)]\(_i\). RVD elicited by a 43% hypsomotic solution, under external and internal Ca\(^{2+}\) chelation, in a fura-2-loaded NG108-15 cell. The cell was incubated with BAPTA/AM (100 μM) for 2 h and kept in a Ca\(^{2+}\)-free isosmotic solution containing EGTA (0.5 mM) for 11 min before the Ca\(^{2+}\)-free 43% hypsomotic challenge. Note that the apparent [Ca\(^{2+}\)]\(_i\), measured as the ratio 358/380, decreases upon exposure to the hypsomotic solution but the RVD response persists.
In seven fura-2-loaded cells treated following the above protocol, upon exposure to the Ca$^{2+}$-free ~40% hyposmotic (42.6 ± 0.4%) solution, the cells swelled to a maximum of 62 ± 4% above their initial volume. The time to reach the maximum swelling was 4.9 ± 0.5 min. After a delay of 54 ± 8 s, RVD ensued at an initial average rate of −1.6 ± 0.5% min$^{-1}$ with 30 ± 6% partial recovery (percent RVD). The mean values of all these variables were smaller than those measured in fura-2-loaded control cells without BAPTA, but the differences were statistically significant only for the initial rate of RVD (Table III and Fig. 9 A). However, the differences in the average rate and extent of RVD in the Ca$^{2+}$-free EGTA-BAPTA-treated cells loaded with fura-2 compared with the calcine-loaded control cells were statistically significant ($P < 0.05$). The Ca$^{2+}$ chelation effects on the extent of RVD responses were not statistically significant when compared with the control cells loaded with fura-2 (Table III), probably because the latter indicator was already exerting some chelating or other side effects in the control group. In other words, the control responses in the fura-2-loaded cells were slightly blunted from the beginning. This can be appreciated by comparing the “control” parameters between calcine-loaded and fura-2-loaded cells (Table I and Fig. 9). Again, although the differences between controls are not statistically significant, the tendencies are obvious.

These results prove that RVD can be produced in the absence of a rise in internal Ca$^{2+}$, suggesting that if a single sequential process underlies RVD, a rise in [Ca$^{2+}$]i cannot be the triggering signal. The effects of Ca$^{2+}$ removal suggest that Ca$^{2+}$ may be acting as a modulator affecting RVD kinetics, or chelation may affect RVD through secondary mechanisms.

**Table III**

<table>
<thead>
<tr>
<th>Variable</th>
<th>Control</th>
<th>Ca$^{2+}$-free</th>
<th>$P$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Osmolality (%)</td>
<td>$n = 10$</td>
<td>−41.4 ± 0.6</td>
<td>NS</td>
</tr>
<tr>
<td>Peak swelling (%)</td>
<td>$n = 7$</td>
<td>−42.6 ± 0.4</td>
<td>NS</td>
</tr>
<tr>
<td>Osmometric</td>
<td>59 ± 5</td>
<td>62 ± 4</td>
<td>NS</td>
</tr>
<tr>
<td>Time to peak (min)</td>
<td>5.0 ± 0.4</td>
<td>4.9 ± 0.5</td>
<td>NS</td>
</tr>
<tr>
<td>RVD latency (s)</td>
<td>90 ± 23</td>
<td>54 ± 8</td>
<td>NS</td>
</tr>
<tr>
<td>RVD rate (% min$^{-1}$)</td>
<td>−5.0 ± 0.5</td>
<td>−1.6 ± 0.5</td>
<td>&lt;0.05</td>
</tr>
<tr>
<td>% RVD</td>
<td>45 ± 8 ($n = 7$)</td>
<td>30 ± 6 ($n = 6$)</td>
<td>NS</td>
</tr>
</tbody>
</table>

Values are means ± SEM.

**The Rate of RVD Responses Elicited with Ca$^{2+}$-free Hyposmotic Challenges Is Not Affected by Restoring External Ca$^{2+}$**

An increase in [Ca$^{2+}$]i in isosmotic media produces shrinkage of N1E-115 cells (Crowe et al., 1995). This suggests that these cells are endowed with Ca$^{2+}$-sensitive mechanisms for net solute efflux, which can be activated by Ca$^{2+}$ in isosmotic media. The question arises as to whether such Ca$^{2+}$-triggered mechanisms can be activated during osmotic swelling. To examine this question, experiments were done in which [Ca$^{2+}$]i was increased during ensuing RVD responses elicited by exposure to a Ca$^{2+}$-free EGTA hyposmotic solution. The rationale was that if Ca$^{2+}$-dependent processes are important, then the rate of cell volume recovery would be enhanced by an increase in [Ca$^{2+}$]. In these experiments, we took advantage of the fact that after superfusion with a Ca$^{2+}$-free solution, restoration of [Ca$^{2+}$]i re-

![Figure 9](https://example.com/figure9.png)

**Figure 9.** Effect of extracellular and intracellular Ca$^{2+}$ chelation on the rate (A) and extent (B) of RVD in cells loaded with calcine or fura-2. The initial rate of RVD (percent at min$^{-1}$) and the extent of RVD (percent RVD at 20 min) were measured as described in Fig. 2. There is a significant difference in the initial average rate of RVD (A), between control and Ca$^{2+}$-chelated (0 Ca$^{2+}$-EGTA/BAPTA) conditions in both calcine-loaded ($P < 0.01$) and fura-2-loaded cells ($P < 0.05$). There is a significant difference ($P < 0.01$) in the extent of RVD (B), between control and Ca$^{2+}$-chelated (0 Ca$^{2+}$-EGTA/BAPTA) conditions in calcine-loaded cells, but not in fura-2-loaded cells.
sults in an increase in $[\text{Ca}^{2+}]$. Fig. 10A shows one such experiment in a fura-2-loaded cell. Note that upon removal of external $\text{Ca}^{2+}$ in the isosmotic solution, the basal level of $[\text{Ca}^{2+}]$, decreased. When a $\text{Ca}^{2+}$-free 40% hypsomotic solution was applied, the relative cell volume increased and the cell started to downregulate its volume. A $\text{Ca}^{2+}$-containing hypsomotic solution was then applied while the cell was still undergoing RVD. This resulted in the expected transient increase in $[\text{Ca}^{2+}]$. However, the rate of RVD (Fig. 10A, top) did not increase. Similar experiments were conducted in calcine-loaded cells ($n = 7$). An example is shown in Fig. 10B in which readmission of external $\text{Ca}^{2+}$ during an RVD response elicited by exposure to a 40% $\text{Ca}^{2+}$-free hypsomotic solution did not alter its time course.

The results obtained in six calcine-loaded cells are summarized in Fig. 11. External $\text{Ca}^{2+}$ was readmitted in a time window between 8 and 14 min after RVD ensued. The time at which external $\text{Ca}^{2+}$ was readmitted in each experiment is indicated by the large gray circles. The filled circles correspond to mean values ± SEM of the six RVD responses. No sign of enhancement of RVD rate upon external $\text{Ca}^{2+}$ readmission was observed.

Other experiments were done ($n = 3$) in which $[\text{Ca}^{2+}]$, was increased during osmotic swelling by adding 1 $\mu$M ionomycin in fura-2-loaded cells. The results were similar; i.e., no significant enhancement of RVD rate could be demonstrated. On the basis of these results, we suggest that $\text{Ca}^{2+}$ is not a triggering signal for RVD in osmotically swollen cells, at least within the explored time window.

**Discussion**

**Calcium-dependent and Calcium-independent RVD**

The nature of the signal transducing the change in cell volume to the activation of osmolyte efflux pathways for RVD in osmotically swollen cells is not clear. Two major models have been proposed, one involving $\text{Ca}^{2+}$ as an intracellular second messenger and the other in which RVD is mediated through $\text{Ca}^{2+}$-independent mechanisms. According to the first model, cell swelling causes an increase in cytosolic $\text{Ca}^{2+}$ that activates volume regulatory mechanisms. The increase in $[\text{Ca}^{2+}]$, may result from $\text{Ca}^{2+}$ influx, via plasma membrane channels (Christensen, 1987), or $\text{Ca}^{2+}$ release from intracellular stores (McCarty and O’Neil, 1992; Wu et al., 1997). The proposed effectors are mostly $\text{Ca}^{2+}$-activated cation and anion channels and carriers. This model has been widely accepted as a paradigm for the involvement of $\text{Ca}^{2+}$ as a transducing signal for RVD. Nevertheless, the existing evidence backing up this model is often weak and fragmentary, and the comprehensive analysis necessary to establish an active role of cytosolic $\text{Ca}^{2+}$ for RVD is lacking in many of the cells in which it has been implicated (Foskett, 1994). One of the experimental facts that is missing in the work claiming a signaling role for $\text{Ca}^{2+}$ in RVD is the simultaneous measurement of changes in $[\text{Ca}^{2+}]$, and cell volume, and the demonstration that the $\text{Ca}^{2+}$ signal is the cause of RVD and not just a parallel phenomenon. In the second model, RVD is effected by $\text{Ca}^{2+}$-independent mechanisms (Cahan and Lewis, 1988; Grinstein and Smith, 1990; Foskett et al., 1994; Lippmann et al., 1995). According to this model, hypsomotic swelling opens anion channels, causing Cl$^{-}$ efflux with consequent cell membrane depolarization towards the equilibrium potential for Cl$^{-}$. This depolarization activates voltage-gated K$^{+}$ channels and increases the driving force for K$^{+}$ loss. The end result is the electrically coupled exit of K$^{+}$ and Cl$^{-}$, accompanied by water. The volume-activated anion chan-
nals may be sufficiently nonselective to permit efflux of organic anions, as well as inorganic anions such as Cl\(^-\) and HCO\(_3\)\(^-\) (Strange et al., 1996). The mechanism whereby these anion channels are activated upon cell swelling remains to be elucidated. However, with the exception of a recent study in human neuroblastoma cells (Basavappa et al., 1995), there is general agreement that Ca\(^{2+}\) is not the activating signal of these anion channels (Doroshenko and Neher, 1992; Strange et al., 1996). In addition to channel-mediated solute efflux, there are carrier-mediated efflux pathways such as electroneutral KCl cotransport that can be activated in swollen cells in the virtual absence of Ca\(^{2+}\) (Thornhill and Laris, 1984; Kramhøft et al., 1986).

Unlike previous reports, in the present study we measured changes in CWV and [Ca\(^{2+}\)], simultaneously in the same cells. We found that in single N1E-115 neuroblastoma cells and in NG108-15 hybrid neuroblastoma × glioma cells, there is a close parallel between osmotic swelling, increase in [Ca\(^{2+}\)], and RVD. The extent and rate of rise of [Ca\(^{2+}\)] and RVD were proportional to the degree of medium hypotonicity and the change in cell volume. Moreover, the correlation between the increase in cell volume, the initiation of the rise of [Ca\(^{2+}\)], and subsequent [Ca\(^{2+}\)], changes, and the onset of RVD show the appropriate time sequence to suggest that a rise in [Ca\(^{2+}\)], might, in principle, mediate RVD. However, cells bathed in Ca\(^{2+}\)-free EGTA solutions, loaded with both the Ca\(^{2+}\) indicator fura-2, showed a robust RVD without a concomitant rise in [Ca\(^{2+}\)]. Moreover, the persistent RVD occurred even when [Ca\(^{2+}\)], decreased below resting levels. Thus, no cause–effect relation exists between the rise in intracellular Ca\(^{2+}\) and the RVD response. These results directly prove for the first time that if a single process underlies RVD, a rise in [Ca\(^{2+}\)], is not the triggering signal. The notion that Ca\(^{2+}\) plays no role in the activation of the normal RVD response has been suggested by other investigators based on less direct evidence from experiments on cell populations in which no simultaneous measurements of intracellular Ca\(^{2+}\) and volume were made (e.g., Jørgensen et al., 1997). Cell populations may or may not be homogeneous and may or may not respond synchronously to osmotic challenges. Therefore, experiments in cell populations do not allow reliable inferences and definitive conclusions about what is happening in single cells, particularly when the relevant variables (changes in cell volume and [Ca\(^{2+}\)]) are not recorded simultaneously.

The rate and extent of the RVD responses that persisted after Ca\(^{2+}\) chelation were reduced with respect to control responses. External and internal Ca\(^{2+}\) chelation may attenuate the RVD response through secondary effects; e.g., membrane depolarization leading to a decreased driving force for Cl\(^-\) efflux or side effects of BAPTA such as intracellular acidification or loss of K\(^+\) and Cl\(^-\) (Jørgensen et al., 1997). Alternatively, the results could be the consequence of the operation of two
parallel processes activated by osmotic swelling, one of which is Ca\textsuperscript{2+} independent. In this model, the rate and extent of RVD would be affected by Ca\textsuperscript{2+} chelation simply by the elimination of the Ca\textsuperscript{2+}-sensitive component. The role of Ca\textsuperscript{2+} in the latter component is not clear at this time. It could be that a certain resting level of intracellular (and extracellular) Ca\textsuperscript{2+} may be necessary for optimal RVD responses, as has been suggested for a medullary thick ascending limb cell line derived from rabbit kidney (Montrose-Rafizadeh and Guggino, 1991). According to this view, Ca\textsuperscript{2+} might play a permissive role, for example acting as a cofactor or maintaining cell integrity, rather than acting as a second messenger.

**Ca\textsuperscript{2+} Regulatory Mechanisms in Anisosmotic and Isosmotic Media**

Our conclusion for a Ca\textsuperscript{2+}-independent RVD is strengthened by the fact that the rate of cell volume recovery was not enhanced by restoring external Ca\textsuperscript{2+} within an interval of 8–14 min after RVD onset. Readmission of external Ca\textsuperscript{2+} always resulted in an increase in [Ca\textsuperscript{2+}], indicating that the lack of effect of external Ca\textsuperscript{2+} readdition on the rate of RVD was not due to block of Ca\textsuperscript{2+} entry during the regulatory response. Similar results were obtained by adding the Ca\textsuperscript{2+} ionophore ionomycin (1 μM) during RVD (not shown). These experiments do not disprove that an increase in [Ca\textsuperscript{2+}], could have a role at earlier times during the RVD response, as has been suggested for cell populations of renal proximal tubules (McCarty and O’ Neil, 1992). Unfortunately, we cannot reintroduce Ca\textsuperscript{2+} at earlier times and have a reliable measurement of the change in RVD rates because there is not sufficient sampling time to assess the change in the slope. The experiment is also not feasible using a two-hyposmotic-challenges protocol because the second response exhibits inactivation. Interestingly, we (Crowe et al., 1995) and others (Hoffmann et al., 1984; Foskett et al., 1994) have shown that elevating [Ca\textsuperscript{2+}], in cells maintained in isosmotic medium produces cell shrinkage. On the basis of these results, we suggest that in addition to the Ca\textsuperscript{2+}-independent volume-regulatory machinery that is activated by cell swelling in hyposmotic media, cells also possess Ca\textsuperscript{2+}-activated transport pathways that result in cell shrinkage in isosmotic media.

**Ca\textsuperscript{2+} Signals in Osmotically Swollen Cells Loaded with Fura-2**

The apparent increase in [Ca\textsuperscript{2+}] upon cell swelling produced by hyposmotic solutions could be due to a change in the $K_d$ of fura-2 for Ca\textsuperscript{2+} that accompanies a decrease in internal ionic strength (Uto et al., 1991), rather than a genuine increase in [Ca\textsuperscript{2+}]. Such a change in internal ionic strength might be expected to accompany a dilution of the internal constituents with the hyposmotic challenge, but we have no independent measure of such changes. We have two arguments to suggest that the monitored Ca\textsuperscript{2+} signals are not spurious. First, we note that they disappear when the cells are pretreated with BAPTA (Fig. 8). This argument is somewhat weakened by the fact that the BAPTA might essentially remove the free internal Ca\textsuperscript{2+} so that the contention that the fura-2 affinity for Ca\textsuperscript{2+} becomes higher is irrelevant since there would not be sufficient [Ca\textsuperscript{2+}], to react with the dye. However, the argument becomes stronger for the case in which external Ca\textsuperscript{2+} was chelated with EGTA. In these cells, as in those loaded with BAPTA, the apparent [Ca\textsuperscript{2+}], does not increase upon osmotic swelling, but actually decreases as a consequence of intracellular dilution (Fig. 10). The second argument is the demonstration that the Ca\textsuperscript{2+} signal is often absent in the second of two identical hyposmotic challenges even though the dilution of the cell contents is the same as in the first hyposmotic challenge, which always results in a transient increase in intracellular Ca\textsuperscript{2+} (Fig. 6). The role, if any, of the rise of [Ca\textsuperscript{2+}], observed during osmotic swelling in the presence of extracellular Ca\textsuperscript{2+} is not clear at this time. This change in [Ca\textsuperscript{2+}], may be a modulating influence or just an epiphenomenon in the RVD process.

**A Signaling Role for Ca\textsuperscript{2+} in RVD?**

Four pieces of evidence are usually given to propose an active role for Ca\textsuperscript{2+} as the signal coupling cell volume expansion to RVD responses (McCarty and O’Neil, 1992; Foskett, 1994). Since these arguments are crucial for the interpretation of the results of the present study, it is worth examining them. First, abolition of RVD when a cell is swollen with a Ca\textsuperscript{2+}-free hyposmotic solution containing chelators is often given as proof of a signaling role for Ca\textsuperscript{2+} (e.g., O’Connor and Kimelberg, 1993). However, chelation in conjunction with the osmotic shock is known to depolarize cells (Berman et al., 1994). In addition, the osmotic challenge results in dilution of cell contents. All these factors, individually or in concert, could alter the driving forces for ion efflux or shift the activation curves for voltage-sensitive channels involved in RVD as a consequence of changes in surface potential. On the other hand, persistence of RVD upon external Ca\textsuperscript{2+} removal does not exclude the possibility of an osmotically induced release of Ca\textsuperscript{2+} from intracellular stores (McCarty and O’Neil, 1992), which could be acting as the second messenger.

The second argument generally offered to propose a signaling role for Ca\textsuperscript{2+} on RVD is the disappearance of the response upon external and internal Ca\textsuperscript{2+} chelation with EGTA and BAPTA, respectively (e.g., Adorante and Cala, 1995). The conclusions from this kind of experiment must also be treated with caution for the
same reasons discussed for external Ca\(^{2+}\) chelation. Conversely, persistence of RVD in cells loaded with a Ca\(^{2+}\) chelator has been interpreted as proof of a Ca\(^{2+}\)-independent RVD (Lippmann et al., 1995). However, unless the adequacy of internal Ca\(^{2+}\) chelation is verified, as was done in the present study, this conclusion is not warranted.

The third piece of evidence often given as proof for Ca\(^{2+}\)-dependent RVD is the block of the response with Ca\(^{2+}\) entry blockers. In particular, these include blockers of voltage-gated Ca\(^{2+}\) channels (e.g., dihydropyridines) or blockers of mechanogated channels (e.g., gadolinium). However, such blockers may have effects on mechanisms other than those involved in Ca\(^{2+}\) entry (Foskett, 1994; Sánchez-Olea et al., 1995; Hamill and McBride, 1996).

The fourth argument usually given in favor of the Ca\(^{2+}\) hypothesis for RVD signaling is that hyposmotic swelling induces an increase in [Ca\(^{2+}\)]. Although a necessary logical condition for postulating a signaling role for Ca\(^{2+}\), it must be proved that the temporal relation between the change in [Ca\(^{2+}\)], and the onset of RVD are appropriate and, more importantly, that the Ca\(^{2+}\) increase is necessary for RVD and not just a parallel phenomenon.

Some investigators (e.g., Hoffmann et al., 1984; Gala, 1986) have suggested that swelling causes a change in the Ca\(^{2+}\) affinity of a putative intracellular element of the RVD machinery, and that therefore a rise in Ca\(^{2+}\) is not necessary for Ca\(^{2+}\) to act as a trigger of the RVD response. First, if there was such a change in the Ca\(^{2+}\) sensitivity of some element of the RVD machinery, then the second messenger or triggering signal would not be Ca\(^{2+}\), but the change in affinity for Ca\(^{2+}\) of this putative element of the RVD machinery. Second, we think that this possibility is unlikely to explain our results because [Ca\(^{2+}\)], decreases below resting levels, yet hyposmotic challenges elicit vigorous RVD responses.

**Inactivation of RVD upon Sequential Hyposmotic Challenges**

We show that the RVD response and the concomitant increase in [Ca\(^{2+}\)], were abolished or significantly attenuated upon the second of two identical hyposmotic challenges applied with an interval of 30–60 min. We demonstrated that the RVD inactivation persists in the nominal absence of external Ca\(^{2+}\) and is independent of changes in internal Ca\(^{2+}\). Moreover, we show that inactivation is independent of the dye used to measure cell volume changes. At least three explanations can be offered for these observations. First, if the RVD is mediated by mechanogated channels, then it is conceivable that these channels adapt and do not recover in time for the second hyposmotic challenge (Hamill and McBride, 1994). If there are mechanogated channels present in neuroblastoma cells that mediate Ca\(^{2+}\) influx in addition to those mediating volume regulatory osmolyte effluxes (Falke and Misler, 1989), then both processes (i.e., the rise in intracellular Ca\(^{2+}\) and RVD) would decrease in parallel without being interdependent. Second if RVD is mediated by the efflux of internal solutes from a finite store, then the apparent inactivation could represent the depletion of that store (Larson and Spring, 1984). Third, inactivation could be due to damage of RVD machinery as a consequence of excessive mechanical stress (Hamill and McBride, 1997). We have recently shown that RVD inactivation can be reversed by loading the cells with taurine before the second hyposmotic challenge. On the basis of this observation, we suggest that inactivation is the consequence of depletion of some internal osmolyte store such as taurine (Brodwick et al., 1997).

In summary, our simultaneous measurements of intracellular Ca\(^{2+}\) and volume in single neuroblastoma cells directly demonstrate that an increase in intracellular Ca\(^{2+}\) is not necessary for triggering RVD. In addition, we demonstrate that the RVD response “inactivates” upon repeated challenges, and that this process is also independent of Ca\(^{2+}\). Our results unify conflicting claims in the literature, based on indirect measurements in cell populations, by showing that: (a) RVD can be independent of a rise in intracellular Ca\(^{2+}\), and (b) there is a component of the response that is sensitive to Ca\(^{2+}\) chelation. The role of Ca\(^{2+}\) in the Ca\(^{2+}\)-dependent component, if any, is not clear at this time. Ca\(^{2+}\) chelation may attenuate RVD through secondary effects. Nevertheless, it is clear that RVD itself can be triggered without a rise of [Ca\(^{2+}\)], in the neuronal cell lines investigated in this paper.

