

NIST Technical Note 1425

***Gases for Electrical Insulation and Arc Interruption:
Possible Present and Future Alternatives to Pure SF₆***

L. G. Christophorou
J. K. Olthoff
D. S. Green

Electricity Division
Electronics and Electrical Engineering Laboratory

and

Process Measurements Division
Chemical Science and Technology Laboratory
National Institute of Standards and Technology
Gaithersburg, MD 20899-0001



November 1997

U.S. Department of Commerce
William M. Daley, Secretary

Technology Administration
Gary R. Bachula, Acting Under Secretary for Technology

National Institute of Standards and Technology
Raymond G. Kammer, Director

National Institute of Standards
and Technology
Technical Note 1425
Natl. Inst. Stand. Technol.
Tech. Note 1425
48 pages (Nov. 1997)
CODEN: NTNOEF

U.S. Government Printing Office
Washington: 1997

For sale by the Superintendent of
Documents
U.S. Government Printing Office
Washington, DC 20402

Bibliographic Information

Abstract

The electric power industry's preferred gaseous dielectric (besides air), sulfur hexafluoride (SF_6), has been shown to be a greenhouse gas. In this report we provide information that is useful in identifying possible replacement gases, in the event that replacement gases are deemed a reasonable approach to reducing the use of SF_6 in high voltage electrical equipment. The report focuses on the properties of SF_6 as a dielectric gas and on the data available for possible alternatives to pure SF_6 (i.e., SF_6 alone). On the basis of published studies and consultation with experts in the field, it was attempted to identify alternative dielectric gases to pure SF_6 for possible immediate or future use in existing or modified electrical equipment. The possible alternative gases are discussed as three separate groups: (i) mixtures of SF_6 and nitrogen for which a large amount of research results are available; (ii) gases and mixtures (e.g., pure N_2 , low concentrations of SF_6 in N_2 , and SF_6 -He mixtures) for which a smaller yet significant amount of data are available; and (iii) potential gases for which little experimental data are available.

Keywords

gaseous dielectrics; gas mixtures; gas recycling; global warming; nitrogen; SF_6 ; SF_6 - N_2 mixtures; sulfur hexafluoride

Ordering

Copies of this document are available from the National Technical Information Service, 5285 Port Royal Road, Springfield, VA 22161, at (800) 553-6847 or (703) 487-4650.

ACKNOWLEDGMENTS

The authors benefitted greatly from the suggestions and comments provided by individuals in industry, academia, and government. Shown below are the individuals who contributed by providing information relevant to the report, or by reviewing and commenting on draft materials. While these individuals influenced the development of this document, the authors are solely responsible for the views expressed in the document. The presence of a reviewer's name here is not intended to suggest endorsement or agreement with all of the views contained in this report.

Christophe Boisseau	Electricité de France, FRANCE
Phil Bolin	Mitsubishi Electric Power Products, Inc., USA
Lowell Brothers	Southern Company Services, Inc., USA
John Brunke	Bonneville Power Administration, USA
Ian Chalmers	University of Strathclyde, U.K.
Alan Cookson	National Institute of Standards and Technology, USA
Steinar Dale	ABB Power T&D Company, Inc., USA
Benjamin Damsky	Electric Power Research Institute, USA
Armin Diessner	Siemens, GERMANY
Elizabeth Dutrow	US Environmental Protection Agency, USA
Lois Ellis	Allied Signal, Inc., USA
Fumihiko Endo	Hitachi, JAPAN
Michael Frechette	Hydro-Quebec, CANADA
Yoshikazu Hoshina	Toshiba, JAPAN
Edmund Kuffel	University of Manitoba, CANADA
Richard LaLumondier	National Electrical Manufacturers Association, USA
Donald Martin	G&W Electric Co., USA
Hugh Morrison	Ontario Hydro, CANADA
Koichiro Nakanishi	Mitsubishi Electric Corporation, JAPAN
C. M. A. Nayar	GEC ALSTHOM, FRANCE
Lutz Niemeyer	ABB Corporate Research, SWITZERLAND
Marshall Pace	University of Tennessee, USA
Reinhold Probst	DILO Company, Inc., USA
Bryan Smith	Cryoquip, Inc., USA
Xavier Waymel	Electricité de France, FRANCE
Howard Withers	Air Products and Chemicals, Inc., USA
Roy Wootton	Formerly at Westinghouse, USA

Gases for Electrical Insulation and Arc Interruption: Possible Present and Future Alternatives to Pure SF₆

Contents

1. Introduction	1
1.1 Sulfur Hexafluoride	2
1.2 Principal Uses of SF ₆ by the Electric Power Industry	2
1.3 Concentrations of SF ₆ in the Environment . . .	3
1.4 SF ₆ is a Potent Greenhouse Gas	4
1.5 SF ₆ Substitutes	5
1.6 Scope of this Report	6
2. Properties of Gaseous Dielectrics	6
2.1 Intrinsic Properties	6
2.1.1 Basic Physical Properties	7
2.1.2 Basic Chemical Properties	8
2.2 Extrinsic Properties	8
2.2.1 Reactions and Byproducts	8
2.2.2 Electrical Discharge and Breakdown . .	8
2.3 Other Requirements for Commercialization . .	9
2.4 Properties of Gaseous Insulators for Specific Industrial Uses	9
2.4.1 Circuit Breakers	9
2.4.2 High Voltage Insulation	9
2.4.2.1 Gas-insulated Transmission Lines	9
2.4.2.2 Gas-insulated Transformers . .	10
3. Required Performance and Testing of Gases . . .	10
4. Possible “Universal-Application” Gas Mixtures . .	11
4.1 Insulation	12
4.1.1 Gas-insulated Transmission Lines . . .	12
4.1.2 Gas-insulated Transformers	15
4.2 Interruption	16
4.3 Gas Handling, Storing, Recycling, and Recovering	22
4.4 Discussion	22
5. Other <i>Promising</i> Gases or Mixtures	23
5.1 High-Pressure N ₂	23
5.2 Low-Concentration SF ₆ -N ₂ Mixtures	25
5.3 SF ₆ -He Mixtures	26
6. Other <i>Possible</i> Substitutes: Future R&D	27
7. Conclusions and Recommendations	28
8. References	29
Appendix A – Sales of SF ₆	35
Appendix B – Issues	37
Appendix C – Potential Barriers	39

1. Introduction

Sulfur hexafluoride (SF₆), the electric power industry's presently preferred gaseous dielectric (besides air), has been shown to be a greenhouse gas. Concerns over its possible impact on the environment have rekindled interest in finding replacement gases. In this report we provide information that is useful in identifying such gases, in the event that replacement gases are deemed a reasonable approach to controlling emissions of SF₆ from high voltage electrical equipment. The report focuses on the properties of SF₆ as a dielectric gas and on the data available for possible alternatives to pure SF₆ (i.e., SF₆ alone). On the basis of published studies and consultation with experts in the field, we attempt to identify alternative dielectric gases to pure SF₆ for possible immediate or future use in existing or modified electrical equipment.

This report first describes the properties that make a good gaseous dielectric, and the tests and measurements that are necessary to demonstrate and document the appropriateness of a gas as a high voltage insulating medium, or for use as an arc or current interrupting medium. An effort has been made to gather expert opinion regarding the possible adoption of likely SF₆ substitutes and the additional tests that are needed to effect their acceptability by electric equipment manufacturers and by the electric power industry. During the preparation of the report, we consulted with a broad spectrum of experts (see Acknowledgments) via a series of meetings on the subject matter and by correspondence. Representatives from electric equipment manufacturers, electric utilities, gas handling and manufacturing companies, and academic institutions were consulted.

An attempt was made during the preparation of this report to identify a gaseous mixture that could be adopted for “universal use” as an immediate replacement of pure SF₆. The large amount of available physical and laboratory data suggest that a 40%SF₆-60%N₂ mixture¹ may exhibit dielectric characteristics suitable for use as insulation in high voltage equipment. However, it is realized that there are difficulties in using this mixture for arc or current interruption, and as a replacement gas in

¹All references in this report to mixtures and concentrations are by volume.

already existing equipment. The reasons for and against the use of this “universal-application” gas mixture are discussed.

The report also discusses other possible substitutes for which a significant but smaller amount of data exists. These include high-pressure pure N₂ and dilute SF₆-N₂ mixtures (concentrations of SF₆ in the mixture less than about 15%) as likely gaseous media for electrical insulation, and SF₆-He mixtures as a possible medium for arc interruption. Other gases and mixtures are also discussed for which the available data are too few to allow an assessment of their utility as a substitute, but which suggest some promise. The need for a future R&D program in these areas is indicated and suggestions are made as to possible elements of such a program. While the literature search utilized in this report was not intended to be complete, it is extensive and can serve as a guide to critical work on alternatives to pure SF₆.

This report concentrates on specific uses of SF₆ by the electric power industry. However, much of the discussion is appropriate for other uses of SF₆ as a high voltage insulating and current interrupting medium.

1.1 Sulfur Hexafluoride

Sulfur hexafluoride is a man-made gas which became commercially available in 1947 [1]. It is one of the most extensively and comprehensively studied polyatomic molecular gases because of its many commercial and research applications.² Its basic physical and chemical properties, behavior in various types of gas discharges, and uses by the electric power industry have been broadly investigated (see, for example, [2–7]).

In its normal state, SF₆ is chemically inert, non-toxic, non-flammable, non-explosive, and thermally stable (it does not decompose in the gas phase at temperatures less than 500 °C). Sulfur hexafluoride exhibits many properties that make it suitable for equipment utilized in the transmission and distribution of electric power. It is a strong electronegative (electron attaching) gas both at room temperature and at temperatures well above ambient, which principally accounts for its high dielectric strength and good arc-interruption properties. The breakdown voltage of SF₆ is nearly three times higher than air at atmospheric pressure [6]. Furthermore, it has good

²Besides the use of SF₆ by the electric power industry, other uses of SF₆ include: semiconductor processing, blanket gas for magnesium casting, reactive gas in aluminum recycling to reduce porosity, thermal and sound insulation, airplane tires, spare tires, “air sole” shoes, scuba diving voice communication, leak checking, atmospheric tracer gas studies, ball inflation, torpedo propeller quieting, wind supersonic channels, and high voltage insulation for many other purposes, such as AWACS radar domes and X-ray machines.

heat transfer properties and it readily reforms itself when dissociated under high gas-pressure conditions in an electrical discharge or an arc (i.e., it has a fast recovery and it is self-healing). Most of its stable decomposition byproducts do not significantly degrade its dielectric strength and are removable by filtering. It produces no polymerization, carbon, or other conductive deposits during arcing, and it is chemically compatible with most solid insulating and conducting materials used in electrical equipment at temperatures up to about 200 °C.

Besides its good insulating and heat transfer properties, SF₆ has a relatively high pressure when contained at room temperature. The pressure required to liquefy SF₆ at 21 °C is about 2100 kPa [5, 8]; its boiling point is reasonably low, –63.8 °C, which allows pressures of 400 kPa to 600 kPa (4 to 6 atmospheres) to be employed in SF₆-insulated equipment. It is easily liquefied under pressure at room temperature allowing for compact storage in gas cylinders. It presents no handling problems, is readily available, and up until recently has been reasonably inexpensive.³ The electric power industry has become familiar and experienced with using SF₆ in electrical equipment.

However, SF₆ has some undesirable properties: it forms highly toxic and corrosive compounds when subjected to electrical discharges (e.g., S₂F₁₀, SOF₂); non-polar contaminants (e.g., air, CF₄) are not easily removed from it; its breakdown voltage is sensitive to water vapor, conducting particles, and conductor surface roughness; and it exhibits non-ideal gas behavior at the lowest temperatures that can be encountered in the environment, i.e., in cold climatic conditions (about –50 °C), SF₆ becomes partially liquefied at normal operating pressures (400 kPa to 500 kPa). Sulfur hexafluoride is also an efficient infrared (IR) absorber and due to its chemical inertness, is not rapidly removed from the earth's atmosphere. Both of these latter properties make SF₆ a potent greenhouse gas, although due to its chemical inertness (and the absence of chlorine or bromine atoms in the SF₆ molecule) it is benign with regard to stratospheric ozone depletion.

1.2 Principal Uses of SF₆ by the Electric Power Industry

Besides atmospheric air, sulfur hexafluoride is the electric power industry's preferred gas for electrical

³From 1960 to 1994 the price of SF₆ in quantity purchases remained basically constant at about \$3.00 per pound (one pound = 0.4536 kg). The current price of SF₆ for quantity purchases in the United States varies from as low as \$12 per lb to over \$37 per pound (\$ 82 / kg) [Private communication, P. Bolin, 1997; P. Irwin, *Electrical World*, February 1997, pp. 27–30].

insulation and for arc quenching and current interruption equipment used in the transmission and distribution of electrical energy. Generally, there are four major types of electrical equipment which use SF₆ for insulation and/or interruption purposes: gas-insulated circuit breakers and current-interruption equipment, gas-insulated transmission lines, gas-insulated transformers, and gas-insulated substations. It is estimated [9–11] that for these applications the electric power industry uses about 80% of the SF₆ produced worldwide, with circuit breaker applications accounting for most of this amount. These estimates are consistent with a recent tabulation of SF₆ production worldwide [12] (See Appendix A). Gas-insulated equipment is now a major component of power transmission and distribution systems all over the world and employs SF₆ almost exclusively. It offers significant savings in land use, is aesthetically acceptable, has relatively low radio and audible noise emissions, and enables substations to be installed in populated areas close to the loads.

Depending on the particular function of the gas-insulated equipment, the gas properties which are the most significant vary. For *circuit breakers* the excellent thermal conductivity and high dielectric strength of SF₆, along with its fast thermal and dielectric recovery (short time constant for increase in resistivity), are the main reasons for its high interruption capability. These properties enable the gas to make a rapid transition between the conducting (arc plasma) and the dielectric state of the arc, and to withstand the rise of the recovery voltage. SF₆-based circuit breakers are presently superior in their performance to alternative systems such as high-pressure air blast or vacuum circuit breakers.

For *gas-insulated transformers* the cooling ability, compatibility with solid materials, and partial discharge characteristics of SF₆ – added to its beneficial dielectric characteristics – make it a desirable medium for use in this type of electrical equipment. The use of SF₆ insulation has distinct advantages over oil insulation, including none of the fire safety problems or environmental problems related to oil, high reliability, flexible layout, little maintenance, long service life, lower noise, better handling, and lighter equipment.

For *gas-insulated transmission lines* the dielectric strength of the gaseous medium under industrial conditions is of paramount importance, especially the behavior of the gaseous dielectric under metallic particle contamination, switching and lightning impulses, and fast transient electrical stresses. The gas must also have a high efficiency for transfer of heat from the conductor to the enclosure and be stable for long periods of time (say, 40 years). SF₆-insulated transmission lines offer distinct advantages: cost effectiveness, high-carrying capacity, low losses, availability at all voltage ratings, no fire risk,

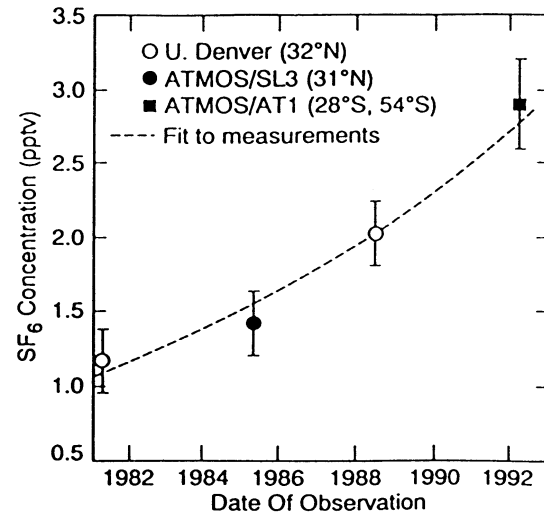


FIG. 1. Average SF₆ concentration (pptv = parts per trillion = parts in 10¹² by volume) between 12 km and 18 km altitude as a function of time [16]. ○ University of Denver balloon-borne infrared measurements at 32° N latitude; ● Spacelab 3 ATMOs data at 31° N latitude; ■ Average of ATMOs ATLAS 1 data at 28° S and 54° S [16].

reliability, and a compact alternative to overhead high-voltage transmission lines in congested areas that avoids public concerns with overhead transmission lines.

Finally, in *gas-insulated substations (GIS)*, the entire substation (circuit breakers, disconnects, grounding switches, busbar, transformers, etc., are interconnected) is insulated with the gaseous dielectric medium and, thus, all of the abovementioned properties of the dielectric gas are significant.

1.3 Concentrations of SF₆ in the Environment

Because of the many and increasing commercial uses of SF₆, there has been an increased demand for it. The estimated world production of SF₆ has increased steadily since the 1970s to approximately 7000 metric tons per year in 1993 [9–11, 13, 14].⁴ In turn, this has resulted in an increased concentration of SF₆ in the atmosphere [11, 13–18]. As seen in Fig. 1, measurements [16, 18] have shown that the amount of SF₆ in the atmosphere has been increasing at a rate of approximately 8.7% per year, from barely measurable quantities a decade ago to current levels near 3.2 pptv (3.2 parts in 10¹² by volume). More recent measurements indicate atmospheric concentrations

⁴This figure is compatible with a compilation of worldwide SF₆ sales data by end-use markets from six companies from the USA, Europe, and Japan (see Appendix A). The total figures listed in Appendix A, however, must be higher than shown, since countries such as China and Russia were excluded from the survey [12].

of SF₆ ranging from 3.18 pptv (at 8 km) to 2.43 pptv (at 27 km) [17a], an atmospheric lifetime of 1937 years [17a], and a global growth rate for atmospheric SF₆ concentrations of 6.9% for 1996 [17b]. While the uncertainties in these numbers make extrapolations difficult, it is clear that the atmospheric concentration of SF₆ is increasing and could reach 10 pptv by the year 2010 [11, 15, 16, 18], depending upon the assumptions of release rates (see Fig. 2).

In some industrial applications SF₆ is not easily recoverable, e.g., in aluminum manufacturing. Releases of SF₆ into the environment by the electric power industry come from normal equipment leakage, maintenance, reclaiming, handling, testing, etc.⁵ Without disposal methods that actually destroy SF₆, it can be expected that all of the SF₆ that has ever been or will ever be produced will eventually enter the atmosphere. This is so even if the present SF₆ leak rate from enclosed power-system equipment is only 1% per year or is improved to < 0.5% per year. It has been suggested [9] that impure, used SF₆ removed from “retired” electrical equipment can be destroyed by thermal decomposition in industrial waste treatment furnaces at elevated temperatures ($T > 1100^{\circ}\text{C}$), but no records are available indicating that this has ever been done at a significant level.

However, decreasing the rate of SF₆ leakage and increasing the level of recycling are high priorities since they both curtail use and production needs of SF₆ and thus reduce the quantities of SF₆ that are eventually released into the environment. Indeed, efforts have recently been undertaken by the electric power industry to reduce and monitor better the amount of SF₆ released into the environment from SF₆-insulated equipment [9–11]. These efforts include:

- *minimizing SF₆ releases* by improved methods to quantify and stop leakages, gradual replacement of older equipment which normally leaks at higher rates, implementation of a sound overall policy of using, handling, and tracing SF₆, better pumping and storage procedures, efficient recycling and setting of standards for recycling [19], and destruction of used SF₆,

- *reducing the amount of SF₆ used* by manufacturing tighter and more compact equipment, development of sealed-for-life electrical apparatus, and replacing SF₆ where possible by other gases or gas mixtures (see later in this report).

These efforts are partially motivated by the prospect

⁵We acknowledge private discussions on these issues with P. Bolin of Mitsubishi Electric Power Products Inc. (USA), J. Brunke of Bonneville Power (USA), H. Morrison of Ontario Hydro (Canada), M. F. Frechette of IREQ (Canada), L. Niemeyer of ABB Research Corporation (Switzerland), A. Diessner of Siemens AG (Germany), K. Nakanishi of Mitsubishi Electric Corporation (Japan), and F. Endo of Hitachi (Japan).

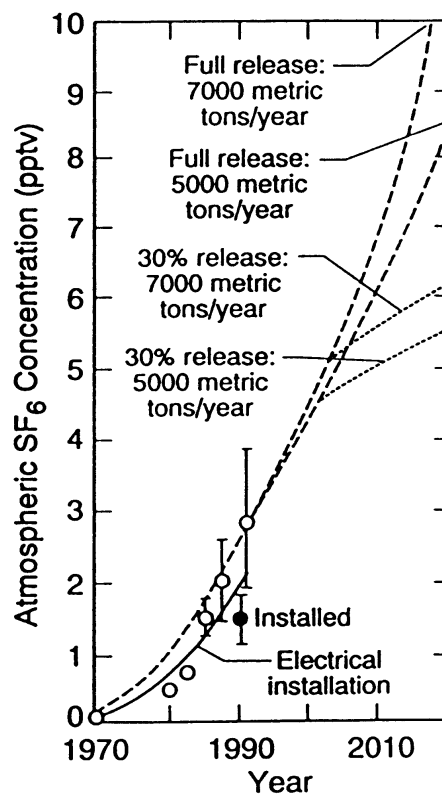


FIG. 2. Atmospheric SF₆ concentration (pptv = parts in 10¹² by volume) as a function of time. The solid curve represents the estimated cumulative total SF₆ from gas-insulated equipment in the past, the open points are measured atmospheric trace concentrations, the solid point labeled “installed” is the estimated concentration assuming that all SF₆ enclosed in electrical equipment throughout the world in 1990 has been released into the atmosphere, and the broken lines are projected increases under various assumptions [11].

of regulation and the possibility of imposition of controls on the use and transport of SF₆ [11, 14, 20] (also see Appendix B for a summary of the current status of regulatory issues related to SF₆ use). The overall concern is motivated by virtually one reason only: *SF₆ is a potent greenhouse gas with an extremely long atmospheric lifetime.*

1.4 SF₆ is a Potent Greenhouse Gas

Greenhouse gases are atmospheric gases which absorb a portion of the infrared radiation emitted by the earth and return it to earth by emitting it back. Potent greenhouse gases have strong infrared absorption in the wavelength range from ~ 7 μm to 13 μm. They occur both naturally in the environment (e. g., H₂O, CO₂, CH₄, N₂O) and as man-made gases that may be released [e. g., SF₆; fully fluorinated compounds (FFC); combustion products such as CO₂, nitrogen, and sulfur oxides]. The effective trapping of long-wavelength infrared radiation

from the earth by the naturally occurring greenhouse gases, and its re-radiation back to earth, results in an increase of the average temperature of the earth's surface. Life on earth depends on a normal greenhouse effect to provide the appropriate temperature for its support. An imbalance in the earth's normal greenhouse effect occurs when the man-made, or anthropogenic, emissions of greenhouse gases contribute to an enhanced greenhouse effect which shifts the balance between incoming and outgoing radiation so that more radiation is retained, thus causing changes in the climate system.

Sulphur hexafluoride is an efficient absorber of infrared radiation, particularly at wavelengths near $10.5\ \mu\text{m}$ [21]. Additionally, unlike most other naturally occurring greenhouse gases (e. g., CO_2 , CH_4), SF_6 is largely immune to chemical and photolytic degradation; therefore its contribution to global warming is expected to be cumulative and virtually permanent. Although the determination of the atmospheric lifetime⁶ of SF_6 in the environment is highly uncertain because of the lack of knowledge concerning the predominant mechanism(s) of its destruction, it is very long; estimates range from 800 years to 3200 years [11, 14, 17, 22–24], with the higher values being the most likely estimates. The strong infrared absorption of SF_6 and its long lifetime in the environment are the reasons for its extremely high global warming potential which for a 100-year time horizon is estimated to be ~24,000 times greater (per unit mass) than that of CO_2 , the predominant contributor to the greenhouse effect [22]. The concern about the presence of SF_6 in the environment derives exclusively from this very high value of its potency as a greenhouse gas.

While the potency of SF_6 as a greenhouse gas is extremely high, the amount of SF_6 in the atmosphere compared to the concentrations of the naturally occurring and other man-made greenhouse gases are extremely low. Estimates of the relative contribution of SF_6 to non-natural global warming – using 1993 estimated SF_6 -concentration levels – range from 0.01% [11] to 0.07% [9, 10]. In 100 years this value could become as high as 0.2% [9]. However, it is feared that SF_6 and other small-volume emissions may have a significant combined influence, and that environmental effects not yet anticipated may be exacerbated by their long lifetime in the atmosphere. Government and environmental protection agencies, electrical, chemical and other industries using or interested in the use of SF_6 [6, 11, 13, 14, 20] have expressed concerns over the possible long-term environmental impact of SF_6 , and the electric power industry is responding in a multiplicity of ways to better control SF_6 usage than in the past and to reduce emissions

into the environment [9–11]. Because SF_6 is already widely used, there are obvious economic implications about any attempts to regulate or control its production, use, and eventual disposal.

Sulfur hexafluoride is an superior dielectric gas for nearly all high voltage applications. It is easy to use, exhibits exceptional insulation and arc-interruption properties, and has proven its performance by many years of use and investigation. However, the extremely high global warming potential of SF_6 mandates that users actively pursue means to minimize releases into the environment, one of which is the use of other gases or gas mixtures in place of SF_6 .

1.5 SF_6 Substitutes

Gaseous insulation must be environmentally acceptable, now and in the future. Therefore, the best response to the concerns over the possible impact of SF_6 on global warming is to prevent the release of SF_6 into the environment. Clearly the most effective way to do this, is not to use SF_6 at all. While such a proposition might be environmentally attractive, it is difficult to envision the near term elimination of the use of SF_6 in view of the industrial reliance on the gas and demonstrated societal value of its use. This environmentally-friendly solution does highlight the need for a search for alternative gaseous insulation and perhaps also the need for alternative high-voltage insulation technologies.

SF_6 -substitute gaseous dielectrics are more difficult to find than it seems on the surface, because of the many basic and practical requirements that a gas must satisfy and the many studies and tests that must be performed to allow confident use. For example, the gas must have a high dielectric strength which requires the gas to be electronegative; however, strongly electronegative gases are usually toxic, chemically reactive, and environmentally damaging, with low vapor pressure, and decomposition products from gas discharges that are extensive and unknown. Non-electronegative gases which are benign and environmentally ideal, such as N_2 , normally have low dielectric strengths. For example, N_2 has a dielectric strength about 3 times lower than SF_6 and lacks the fundamental properties necessary for use by itself in circuit breakers. Nonetheless, such environmentally friendly gases might be used by themselves at higher pressures, or at comparatively lower

⁶The time taken for a given quantity of SF_6 released into the atmosphere to be reduced via natural processes to ~37% of the original quantity.

pressures as the main component in mixtures with electronegative gases, including SF₆, at partial concentrations of a few percent.

The search for SF₆ substitutes traces back many years. It was especially intense in the 1970s and 1980s when gases “superior” to SF₆ were being sought. A number of studies conducted mainly during this time period, produced a large body of valuable information (see, for example, Refs. 2 and 3) which needs to be revisited and be reassessed not from the perspective of finding “better” gaseous dielectrics than SF₆, but rather from the point of view of finding gases or gas mixtures which are environmentally acceptable and comparable in dielectric properties and performance to SF₆. A rekindled interest in “new” gaseous insulators may also direct itself toward finding gases or gas mixtures which are not necessarily universally optimum for every high-voltage insulation need, but which can be optimized for a particular application.

Any program on substitutes needs to address comprehensively the issues involved and evaluate possible substitutes within the framework of the total environment. Besides the obvious requirements of high gas pressure, non-toxicity, non-flammability, availability and cost, there should be basic, applied, and industrial testing to assess the thermal and electrical properties of other gaseous dielectrics. Performance under various voltages (DC, AC, impulse, transients), field configurations, and particle contamination must be tested. Gas decomposition under prolonged electrical stress, corona, breakdown, and arc must be investigated, along with gas aging and the influence of spacer and other materials. Gas mixtures in particular need to be looked at anew. Efforts must be made to address concerns regarding mixtures which include difficulties in handling, mixing, maintaining constant mixture composition, reclaiming of mixture's constituents, possible inferior performance with regard to thermal, insulation, and current interruption properties, and the associated equipment design changes that such use may entail. It must be emphasized, however, that gas mixtures should be tested under conditions (e.g., pressures, equipment design) where they are likely to perform well, not simply under conditions for which SF₆ is better. It must also be stressed that historically resources have not been as abundant for the study of gas mixtures as they had been for the study of pure SF₆.

1.6 Scope of this Report

It is the purpose of this report to provide information regarding the following:

1. The required or desirable properties of any dielectric gas for use in the various applications

of the electric power industry.

2. The tests required to document the suitability and acceptability of a dielectric gas for the intended application(s).
3. The feasibility of a “universal” gas mixture that could substitute for pure SF₆ and help reduce the current levels of SF₆ utilized by the electric power industry.
4. Alternate gases or gas mixtures for which a significant amount of data are available supporting their possible use in newly-designed industrial equipment.
5. Possible gases or gas mixtures for which little physical data are presently available, but which are sufficiently promising to justify further research.
6. Recommendations on substitutes and future R&D aimed at the development of environmentally acceptable alternatives to pure SF₆.

Items 1 to 6 are respectively discussed in Sections 2 to 7 of this report.

2. Properties of Gaseous Dielectrics

The properties of a gas that are necessary for its use in high voltage equipment are many and vary depending on the particular application of the gas and the equipment. They are also interconnected and coupled. In their optimum combination one may achieve distinctly desirable synergisms with regard to dielectric strength, for instance, which clearly show that a gas mixture may be more than just the partial-pressure-weighted addition of the dielectric strength of the individual mixture components [2, 3, 25]. In the following sections, the gaseous dielectric properties which are of particular importance in high voltage applications are identified.

For the purpose of this report the properties of a gaseous dielectric are divided into four groups:

- intrinsic properties (physical and chemical);
- extrinsic properties (reactions, gas byproducts, discharge and breakdown);
- other requirements for commercial use; and
- specific properties required for arc interruption, transmission lines, and transformers.

2.1 Intrinsic Properties

Intrinsic properties are those properties of a gas which are inherent in the physical atomic or molecular structure of the gas. These properties are independent of the application or the environment in which a gas is placed.

2.1.1 Basic Physical Properties

One of the desirable properties of a gaseous dielectric is *high dielectric strength* (higher, for instance, than air). The gas properties that are principally responsible for high dielectric strength are those that reduce the number of electrons which are present in an electrically-stressed dielectric gas. To effect such a reduction in the electron number densities, a gas should:

- be electronegative (remove electrons by attachment over as wide an energy range as possible); it should preferably exhibit increased electron attachment with increasing electron energy and gas temperature since electrons have a broad range of energies and the gas temperature in many applications is higher than ambient;
 - have good electron slowing-down properties (slow electrons down so that they can be captured efficiently at lower energies and be prevented from generating more electrons by electron impact ionization); and
 - have low ionization cross section and high ionization onset (prevent ionization by electron impact).
- The significance of these parameters, especially electron attachment, in determining the dielectric strength of the

TABLE 1. Relative DC uniform-field breakdown strengths V_s^R of some dielectric gases.^a

Gas	V_s^R b,c	Comments
SF ₆	1	Most common dielectric gas to date besides air
C ₃ F ₈	0.90	Strongly and very strongly electron attaching gases, especially at low electron energies
n-C ₄ F ₁₀	1.31	
c-C ₄ F ₈	~1.35	
1,3-C ₄ F ₆	~1.50	
c-C ₄ F ₆	~1.70	
2-C ₄ F ₈	~1.75	
2-C ₄ F ₆	~2.3	
c-C ₆ F ₁₂	~2.4	
CHF ₃	0.27	Weakly electron attaching; some (CO, N ₂ O) are effective in slowing down electrons
CO ₂	0.30	
CF ₄	0.39	
CO	0.40	
N ₂ O	0.44	
Air	~0.30	
H ₂	0.18	
N ₂	0.36	Non-electron attaching but efficient in slowing down electrons
Ne	0.006	Non-electron attaching and not efficient in slowing down electrons
Ar	0.07	

^a Based on Table 2 of Ref. 25.

^b Some of the values given are for quasi-uniform fields and may thus be somewhat lower than their uniform-field values.

^c The relative values listed in the table can be put on an absolute scale by multiplying by 3.61×10^{15} V cm², the uniform-field breakdown field, $(E/N)_{lim}$, of SF₆.

gaseous medium can be seen from the representative data for different gases in Table 1. It is evident in this table that some gases actually exceed the dielectric strength of SF₆. However, they all exhibit negative properties as to make them less desirable gaseous insulators in practical systems as presently designed. Figure 3 illustrates the basic physical properties of electron attachment, ionization, and scattering as they relate to the dielectric strength [25]. The most critical property of a gaseous dielectric for high dielectric strength is a large electron attachment cross section over a wide electron energy range. The second most significant property is a large electron scattering cross section at low electron energies to slow electrons down so that they can be captured more efficiently and be prevented from generating more electrons in collisions with the dielectric gas molecules.

Furthermore, the gas properties must be such that electron detachment from negative ions is prevented since electron detachment is a major source of electrons that trigger gas breakdown. The negative ions that are formed (through the formation of negative ions by electron attachment) must be as stable as possible. Detachment of electrons from negative ions can occur via a number of processes, foremost by autodetachment, collisional detachment, and photodetachment. Especially the former process is a strong function of gas temperature [26].

The measurements needed to quantify the intrinsic physical properties of a gaseous dielectric for insulation include:

- electron attachment cross sections;
- electron scattering cross sections;
- electron impact ionization cross sections;
- electron detachment cross sections (photodetachment, collisional detachment, and the associated processes of clustering and ion-molecule reactions); and
- coefficients for electron attachment, ionization, effective ionization, and transport.

Besides the above properties, there are a number of other basic properties which are necessary for the complete characterization of the dielectric gas behavior and its performance in practice. These include:

- secondary processes such as electron emission from surfaces by ion and photon impact;
- photoprocesses;
- absorption of photoionizing radiation (this is a controlling factor in discharge development in non-uniform fields);
- dissociation under electron impact decomposition;
- ion-molecule reactions;
- reactions with trace impurities; and
- reactions with surfaces.

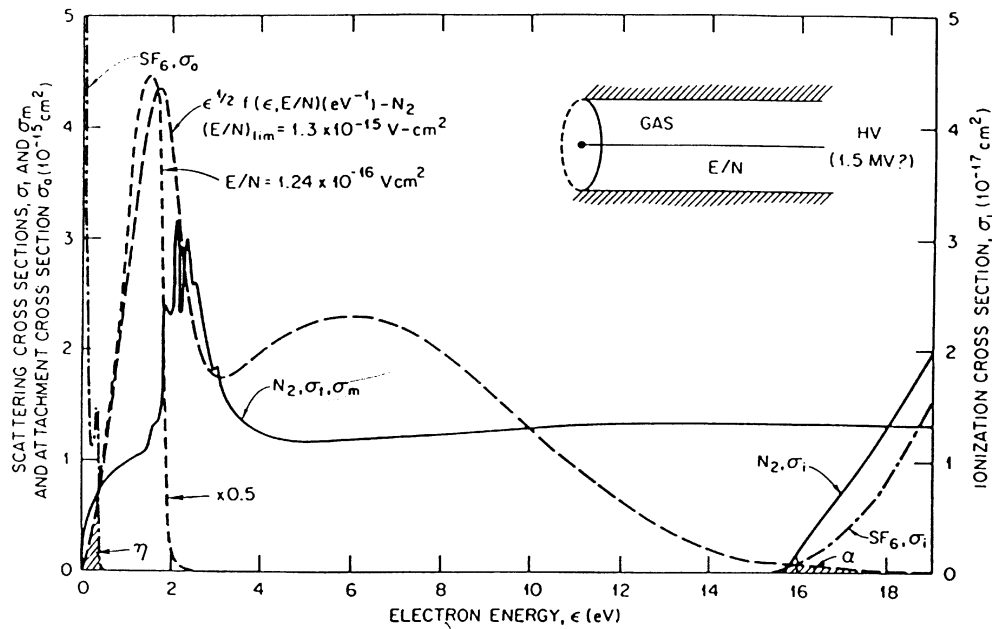


FIG. 3. Total ionization cross section $\sigma_i(\mathcal{E})$ for N_2 (—) and SF_6 (-·-) close to the ionization onset. Total electron scattering cross section $\sigma_s(\mathcal{E})$ as a function of electron energy, \mathcal{E} , for N_2 (—), and total electron attachment cross section $\sigma_a(\mathcal{E})$ for SF_6 (-·-). Electron energy distribution functions in pure N_2 for two values of the density, N , reduced electric field E/N : at a value of $1.24 \times 10^{-16} \text{ V cm}^{-2}$, about ten times lower than the E/N value at which breakdown occurs under a uniform electric field, and at the limiting value of $E/N (= 1.3 \times 10^{-15} \text{ V cm}^{-2})$ at which breakdown occurs under a uniform electric field. The shaded areas designated by η and α are, respectively, a measure of the electron attachment and electron impact ionization coefficients for SF_6 (from [25]).

2.1.2 Basic Chemical Properties

The dielectric gas must have the following “chemical” properties:

- high vapor pressure;
- high specific heat (high thermal conductivity) for gas cooling;
- thermal stability over long periods of time for temperatures greater than 400 K;
- chemical stability and inertness with regard to conducting and insulating materials;
- non-flammable;
- non-toxic; and
- non-explosive.

When used in mixtures, it must have appropriate thermodynamic properties for mixture uniformity, composition, and separation (see Appendix C).

2.2 Extrinsic Properties

Extrinsic properties are those which describe how a gas may interact with its surroundings, or in response to external influences, such as electrical breakdown and discharges.

2.2.1 Reactions and Byproducts

To be used in electrical applications, a dielectric gas should:

- undergo no extensive decomposition;
- lead to no polymerization;
- form no carbon or other deposits; and
- be non-corrosive and non-reactive to metals, insulators, spacers, and seals.

In addition it should have:

- no major toxic or adversely-reactive byproducts;
- removable byproducts; and
- a high recombination rate for reforming itself, especially for arc interruption.

Finally, the gas must be environmentally friendly, e.g., it must not contribute to global warming, must not deplete stratospheric ozone, and must not persist in the environment for long periods of time.

2.2.2 Electrical Discharge and Breakdown Properties

Specific properties of the gas under discharge and breakdown conditions include:

- a high breakdown voltage under uniform and

- non-uniform electric fields;
- insensitivity to surface roughness or defects and freely moving conducting particles;⁷
- good insulation properties under practical conditions;
- good insulator flashover characteristics
- good heat transfer characteristics;
- good recovery (rate of voltage recovery) and self-healing;
- no adverse reactions with moisture and common impurities; and
- no adverse effects on equipment, especially on spacers and electrode surfaces.

Also some knowledge must be available concerning its discharge mechanisms (corona, breakdown, arc) and discharge characteristic behavior, and its decomposition under arc and various types of discharges.

2.3 Other Requirements for Commercialization

Commercial use of a dielectric gas requires certain non-physical characteristics, including widespread availability, reliable supply, and long-range stability of supply.

2.4 Properties of Gaseous Insulators for Specific Industrial Uses

2.4.1 Circuit Breakers – Arc Quenching and Current Interruption

An electric arc is the most crucial switching element in a circuit breaker. It has the unique ability to act as a rapidly changing resistor such that during the AC current, high conductance is maintained. As the current approaches zero, the conductance decreases rapidly, and finally, at zero current, the resistance rises to prevent re-ignition. Commercial circuit breakers utilize air, oil, SF₆, solid state, or vacuum as interrupting media. The arc properties for gas-based circuit breakers are a strong function of the arcing gaseous medium. The most significant required gas properties for arc interruption are:

- *High dielectric strength comparable to that of sulfur hexafluoride* - This is one of the most essential properties characterizing a good interrupting medium.
- *High thermal conductivity* - This is another

⁷The optimum design of a gas-insulated system requires this knowledge. Perhaps one can determine this through the so-called figure of merit, i.e., from basic measurements of $(\alpha - \eta)/N$ versus E/N . It would certainly be desirable to have a gas for which these effects are less troublesome than for SF₆.

important required property. The arc is initially hot (temperatures in excess of 10,000 K), and it must be quickly cooled down by removing energy from it by the gas. Additionally, the arc must have a short time constant for the increase in resistivity. For these requirements, the gas must have high thermal conductivity at high temperatures and also should capture quickly free electrons when the gas is hot and the electrons fast. These two properties – high thermal conductivity and high electron attachment – lead to a high interruption capability, i.e., enable a rapid transition between the conducting state (arc plasma) and a dielectric state able to withstand the rise of recovery voltage. SF₆ is known to have a time constant 100 times shorter than air and is used in circuit breakers for two main reasons: it has a high thermal conductivity at high temperatures which enables it to rapidly cool down; SF₆ and its decomposition products are electronegative and thus enhance the disappearance of electrons even when the gas is hot.

- *Fast gas recovery* - At the high temperatures involved, the gas molecules are dissociated into their constituent atoms (atomized). They must quickly reassemble, preferably to form their original molecular structure. (Besides SF₆, this is a property shared by a number of molecules with top symmetry such as σ -bonded perfluoroalkanes).

- *Self-healing / dielectric integrity* - This limits the preferred gases to those that are either atomic in nature or molecular with very compact and stable structure, such as SF₆, CF₄, and other compounds, which when “atomized” under the high temperature arc conditions reform themselves with high efficiency, that is, the original molecules are the main decomposition product.

2.4.2 High Voltage Insulation

There are two important types of basic gas-insulated apparatus used by the electric power industry: gas-insulated transmission lines and gas-insulated transformers. In this section are outlined some of the principal properties a gaseous dielectric needs to be used in such applications. Other applications with similar needs include buses and disconnects in gas-insulated substations.

2.4.2.1 Gas-insulated Transmission Lines - Here the dielectric strength of the gas and its long-range stability and inertness, along with its heat transfer properties at temperatures much lower than in circuit breakers (≤ 110 °C), are important gas requirements. Specifically, the required properties include:

- high dielectric strength (in uniform fields, non-uniform fields, in the presence of electrode

- roughness and conducting particles, and for various geometries including co-axial configurations);
- high vapor pressure at operating and ambient temperature;
- chemical inertness;
- high thermal conductivity [but at temperatures far below those encountered in arcs (a few hundred degrees above ambient)];
- no thermal aging (long-term, 40 years or more);
- no deposits (no carbon deposits, no polymerization, and no decomposition);
- easily removable, non-harmful byproducts;
- no hazards (fire, explosion, toxicity, corrosion).

2.4.2.2 Gas-insulated Transformers - In very early transformers, air was the most commonly used insulating medium, but as the voltages were increased, oil was substituted for air. While oil is presently widely used and has many advantages, it burns when exposed to flame or heated to ignition point in the presence of air. Also, certain mixtures of oil vapor and air explode on ignition when confined. Additionally, breakdown due to charge accumulation on insulating parts by ions transported by the cooling pumps may occur, and flashovers due to particulate contaminants may be caused.

There are distinct advantages in using gas insulation in transformers. Firstly, the use of a gas instead of oil completely removes the undesirable characteristics of oil just mentioned. Secondly, gas-filled transformers are lighter, have better noise characteristics (since gas transmits less vibration than oil), and are easier to handle. Compared to oil, however, the gas is not as good for cooling (needs special techniques to remove the heat) and thus gas-insulated transformers presently are unable to meet the highest ratings achieved by oil transformers. The properties of the gas required for this application include:

- high dielectric strength at reasonable (e. g., 500 kPa) pressures;
- low boiling point (low condensation temperature, high vapor pressure);
- low toxicity;
- chemical inertness;
- good thermal stability (because transformers are operated in a wide temperature range);
- non-flammable;
- high cooling capability (heat transfer is important in transformers which frequently get quite hot);
- good compatibility with solid materials (because the gas must coexist with many different solid materials in the gas-insulated transformer);
- good partial discharge characteristics (because of the high possibility of partial discharges in the transformer);

- useable over a range of temperatures (basic properties as a function of temperature);
- safe, easy to handle, inexpensive, securely available.

3. Required Performance and Testing of Gases

At first consideration one may be tempted to adopt an extreme position for new gases: *ALL that has been done on SF₆ has to be repeated*. While there is a need for any new gas “to be proven,” this approach is unrealistic, impractical, and perhaps unwise and unnecessary. Clearly, before any testing is done, the gas must:

- be environmentally acceptable, or confined for life,
- have no serious known health-related risks and serious safety-related problems (toxicity, flammability, etc.),
- have a high pressure (to be useful as a unitary gas or as an additive in mixtures), and
- be available, stable, and thermally and chemically inert.

These requirements must be satisfied whether one is looking for potential gas substitutes on which tests have already been made or for new gaseous systems for which tests will be made, independently of the intended use.

The list of other tests that are also useful and desirable is long (see Sec. 2 on required properties) and includes:

- breakdown tests as a function of pressure, field, types of voltage, and time;
- comprehensive dielectric strength tests using practical-size systems and voltages and waveforms (i.e., DC, AC, lightning and switching impulse, fast transients). Since the design of the high voltage insulation system is usually determined by the lightning impulse test level (BIL), the lightning impulse test is a crucial test;
- effects of surface roughness and conducting particles. Practical design levels for the dielectric strength are normally much lower than the “theoretical” dielectric strength of a gas insulator, because the dielectric strength of gases, especially those for strongly electronegative gases, are very sensitive to field perturbations such as those caused by conductor surface imperfections and by conducting particle contaminants;⁸
- dielectric strength measurements at high gas

⁸The design levels for SF₆ have been quoted [27] to be of the order of 37% of the theoretical strength of SF₆ for lightning impulse and 19% of the 60 Hz factory test for these stated reasons.

pressures (this is one type of measurement that has generally been lacking and is crucial);

- long-time tests;
- flashover voltages of insulators;
- thermal stability in the presence of other materials (long-time stability with metals and resins), and thermal aging;
- corona inception⁹ and extinction
- thermal cooling.
- mass and light spectroscopy to identify the discharge products and their reactions for a number of purposes including diagnostics;
- measurement of dielectric strength as a function of gas pressure, especially for weakly electron attaching gases or mixtures;
- scaling data on small laboratory equipment to large practical systems, and extrapolating data taken over short time scales to the expected long life times of industrial systems (e.g., 40 years);
- byproducts and possible health effects.

The list of desirable tests for use of a gas under arc or current interruption applications must also include:

- tests of arc and current interruption properties;
- recovery tests; and
- nozzle design and behavior.

4. Possible “Universal-Application” Gas Mixtures

The most desirable SF₆ substitute would be a gas that could be put in all existing SF₆-equipment, requiring little or no change in hardware, procedures or ratings. Such a gas we refer to as a “universal-application” gas and we define it as a gaseous medium which can be used instead of pure SF₆ in existing equipment without significant changes in practice, operation, or ratings of the existing gas-insulated apparatus. It is a useful exercise to determine if such a substitute can be identified from the existing gaseous dielectric data.

Of the many unitary, binary, and tertiary gases or gas mixtures that have been tested over the last three decades or so, SF₆-N₂ mixtures seem to be the most thoroughly characterized [yet not completely tested, especially at high

⁹It has been pointed out by Wootton [28] that in tests on a full size GIS with a fixed particle, typically less than 10% of the breakdowns occur without corona stabilization. Based on this information, Dale et al. [27] suggested that in practical apparatus it would be the corona inception level and not the corona stabilized breakdown level which is important. However, the strong corona stabilization characteristics of electronegative gases can be advantageous.

pressures (greater than 0.5 MPa)] gaseous dielectric media besides pure SF₆ [2–4, 6–8, 14, 25, 29–34]. There is broad acceptance of the view that these mixtures may be good replacements of pure SF₆. The main reasons are:

- they perform rather well for both electrical insulation applications and in arc or current interruption equipment,
- they have lower dew points and certain advantages especially under non-uniform fields¹⁰ over pure SF₆,
- they are much cheaper than SF₆ especially after the recent large increases² in the price of SF₆, and
- industry has some experience with their use.

The relevant question, therefore, is: *does an optimum mixture composition and total pressure exist that allows the use of this mixture as a “universal-application” gas, and could the industry readily use such a mixture?* While the answer to this question is complex, it is desirable to attempt to identify, on the basis of existing knowledge, a particular mixture composition that may be best suited for consideration by the electric power industry for their needs. If such mixture can be identified, it can perhaps be standardized in composition. Although it would be desirable to have such a standard mixture prepared and sold by chemical companies for direct use in the field, this may not be feasible, and the two gases would probably have to be mixed to the standard composition at the point of use (see Appendix C).

Based upon research conducted world-wide over the last three decades or so, it appears that the optimum composition of an SF₆-N₂ mixture for use by the electric power industry in place of pure SF₆ for both high voltage insulation (for gas-insulated transmission lines and gas-insulated transformers) and arc or current interruption purposes may be in the range of 40% to 50%SF₆ in N₂. Thus, possible standard mixtures that can reasonably be considered are 40%SF₆-60%N₂ or 50%SF₆-50%N₂.

The savings¹¹ of replacing pure SF₆ by a 40%SF₆-60%N₂ gas mixture are potentially large. If it is assumed that 80% of the ~8,000 metric tons of SF₆ produced annually is used by the electric power industry (Sec.1.2; [12]), at a price¹² of \$20 / lb (~\$42 / kg) for SF₆, the total annual savings in the cost of SF₆ will be about \$150 million.

¹⁰The more electronegative the gas is, the larger the reduction of its dielectric strength under non-uniform field conditions and in the presence of conducting particles.

¹¹Perhaps even higher savings may be possible if the percentage of SF₆ is lowered further by increasing the total operating pressure of the mixture. Limited measurements on the arc interruption capability of pure SF₆ in the pressure range 0.41 MPa to 0.72 MPa ([3], p. 51) indicate that it increases almost as the square of the fill pressure.

¹²Based on the spectrum of prices, it seems logical to assume a price of about \$20 per lb (\$42 / kg).

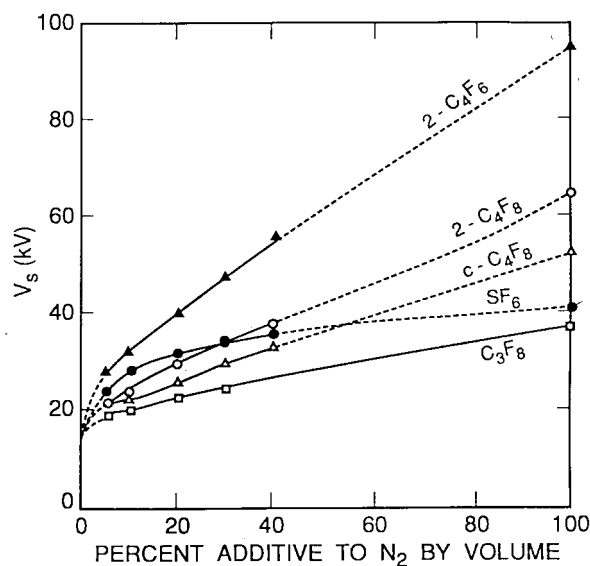


FIG. 4. DC breakdown voltage as function of electron attaching additive to nitrogen (uniform and quasi-uniform electric fields; the total gas pressure is 66.66 kPa and the electrode gap is 7 mm). The broken curves are extrapolations (from [35, 36]).

The feasibility analysis of a universal application mixture for insulation applications, and arc and current interruption purposes is based on information obtained from a number of sources, the most significant of which are briefly discussed below. Additional pertinent information related to mixtures with even lower concentrations of SF_6 is presented in Sec. 5.2.

4.1 Insulation

4.1.1 Gas-insulated Transmission Lines

Existing information suggesting the use of SF_6 - N_2 mixtures for insulation purposes has been summarized and discussed in a number of recent publications (see, for example, Refs. [3, 6, 14, 33, 34]). In this section we refer to and supplement the work summarized in those reports, which indicates the possibility of adopting the 50% SF_6 -50% N_2 mixture as a standard gas option for gas-insulated transmission lines. Most of the information presented also supports the use of such mixtures for gas-insulated transformers (Sec. 4.1.2) and possibly also circuit breakers (Sec. 4.2). It has been known for a long time that the breakdown voltage of SF_6 - N_2 mixtures saturates as the percentage of SF_6 in the binary mixture is increased above about 40%. This is seen from DC measurements ([35, 36], Fig. 4), AC measurements ([37], Fig. 5), and impulse measurements ([37], Fig. 6). Above this saturation level, addition of more SF_6 to N_2 yields limited returns for insulation applications. This has been shown by many

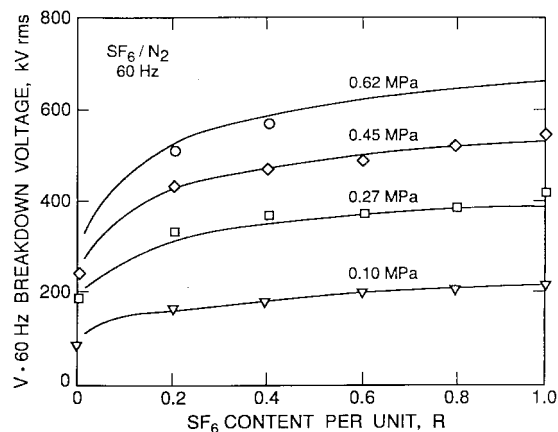


FIG. 5. Measured (symbols) and calculated (solid lines) 60-Hz breakdown voltage values for SF_6 - N_2 in coaxial electrode geometry (from [37]).

studies. For instance, Bouldin et al. [30, 31] conducted an assessment of the potential of dielectric gas mixtures for industrial applications based mostly on DC uniform and quasi-uniform field data at pressures less than a few hundred kPa. They concluded that a 50% SF_6 -50% N_2 mixture, operated at 15% higher pressures, exhibits the same dielectric strength as 100% SF_6 , but at ~35% lower cost (calculated using a price of SF_6 more than ten times lower than it is today). A 50% SF_6 -50% N_2 mixture was listed as having 0.88 the dielectric strength of pure SF_6 at the same pressure and a condensation point 20 °C lower.

Malik and Qureshi [38] reviewed the work on electrical breakdown in mixtures of SF_6 and other gases

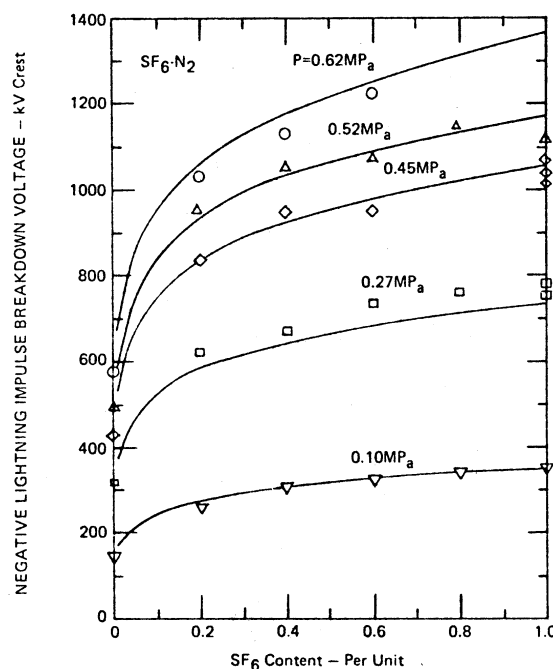


FIG. 6. Negative lightning impulse breakdown voltages for SF_6 - N_2 mixtures. The solid curves are semiempirical fits to the measurements (from [37]).

including N_2 . They state: “ SF_6 - N_2 mixtures containing 50% to 60% of SF_6 have dielectric strength of up to 85% to 90% that of pure SF_6 . Such mixtures can have improved impulse and power-frequency breakdown strength in highly non-uniform field gaps and therefore are expected to be less sensitive (than pure SF_6) to the presence of free conducting particles and electrode surface roughness.” Furthermore, the mixtures can be operated at pressures considerably higher than 600 kPa which is the upper limit for SF_6 -insulated apparatus [39] and at considerable reduction in cost [38].

In view of the superior insulation properties of N_2 at higher pressures and in non-uniform fields compared to pure SF_6 , the advantage of the 50-50 mixture over pure SF_6 may be even greater. For example, non-uniform (positive point-plane electrode geometry) DC breakdown measurements [36] as a function of the total pressure showed that the dielectric strength of a 30% SF_6 -70% N_2 mixture at ~600 kPa is somewhat higher than that of pure SF_6 under the same conditions. Similarly, Nakanishi [40] concluded that “a gas mixture of SF_6 - N_2 (and SF_6 -air) is thought to be the most promising candidate for application to high power apparatus. SF_6 - N_2 (and SF_6 -air) mixtures have breakdown properties superior to pure SF_6 at non-uniform fields.”

A recent example [41] of the voltage versus pressure characteristic of SF_6 and a 75% SF_6 -25% N_2 mixture is shown in Fig. 7. Included in the figure are measurements of both the breakdown and the corona onset voltages for the two gases. The measurements were made using nonuniform fields and lightning impulse voltage. The corona onset for SF_6 and the SF_6 - N_2 mixture scales with the respective uniform field breakdown voltages of the two gases, but the breakdown voltage exhibits the usual corona stabilization region which varies with mixture composition. Depending on the mixture composition there are total pressures for which the breakdown voltage of the mixture exceeds that of pure SF_6 .

Similar conclusions have been reached by other impulse breakdown studies. Lightning and switching impulse breakdown measurements in the pressure range between 0.1 MPa and 0.7 MPa by Cookson and Pedersen [37] led them to conclude that “the SF_6 mixtures with N_2 (or air, or CO_2) look promising for compressed gas-insulated transmission (CGIT) applications with a 50-50 mixture at a typical pressure of 0.54 MPa being able to replace SF_6 at 0.45 MPa without loss of breakdown strength.” Similarly, Cookson [42] concluded that mixtures of SF_6 with N_2 can be readily applied in practical CGIT lines at a cost savings over SF_6 . Rein and Kulsetås [43] studied lightning and switching impulse breakdown of SF_6 - N_2 mixtures using electrode configurations representative of open disconnectors and earthing switches and concentric cylinder systems, and pressures

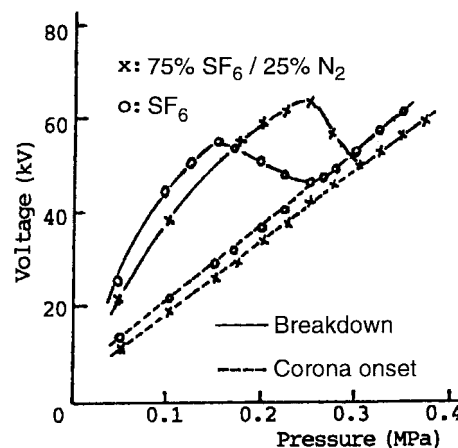


FIG. 7. Voltage versus pressure characteristic of SF_6 and a 75% SF_6 -25% N_2 mixture for a point-plane gap and lightning impulse (+1.2 / 50 μ s) (from Qiu and Feng [41]).

ranging from 0.5 MPa to 1.5 MPa. They concluded that “compared to pure SF_6 of 1.0 MPa, a 50%-50% mixture of SF_6 and N_2 with the same total pressure retains about 85% of the dielectric strength. An increase of N_2 content to a total pressure of 1.5 MPa gives a dielectric strength equal to that of 1.0 MPa SF_6 . The condensation temperature is thereby lowered from $-15^\circ C$ to $-40^\circ C$.”

Furthermore, Fatheddin et al. [44] investigated the breakdown properties of SF_6 - N_2 mixtures using lightning and switching impulses of both polarities, a point-plane geometry, and total pressures between 0.05 MPa and 0.5 MPa. From Figs. 1 to 5 of their paper it appears that a 40% SF_6 -60% N_2 mixture is an excellent choice. For this mixture, the impulse polarity had negligible effect with lightning surges. However, the negative polarity V_{50} (V_{50} is the peak voltage with 50% breakdown probability) values were higher and the positive V_{50} values were lower for the (slower) switching surge pulse. Interestingly, their positive DC point-plane geometry measurements (Figs. 6 and 7B of their paper) showed a 30% SF_6 -70% N_2 mixture to be at least as good as pure SF_6 for total pressure greater than 500 kPa.

Very importantly, a number of studies [3, 29, 45–52] have shown that in the presence of particles a number of SF_6 - N_2 mixtures perform well compared to pure SF_6 . They have also indicated that the effect of particles on the breakdown strength of such mixtures depends on the total pressure and on the partial pressure of the component gases (Fig. 8; Table 2). The data in Table 2 are particularly interesting. They show that for certain conditions (DC, cylindrical electrodes, in the presence of conducting particles as contaminants) the highest breakdown voltage for a 50% SF_6 -50% N_2 mixture corresponds to a total pressure of about 608 kPa (~ 6 atm) (see Table 2), which is a reasonable pressure for use in

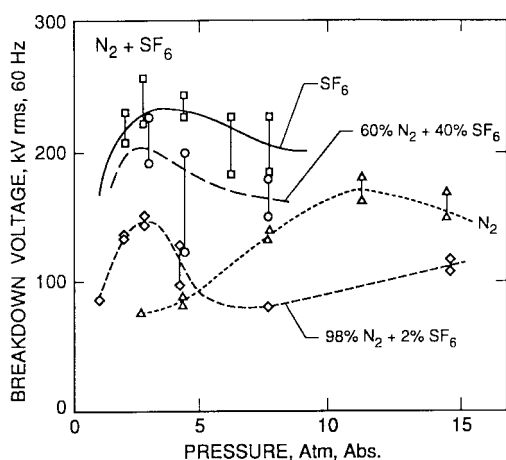


FIG. 8. AC breakdown voltage in SF₆-N₂ mixtures with 0.043 cm (0.017 in) diameter copper wires, 0.635 cm (0.25 in) long in a 7.62 cm (3 in) plane gap [3, 45, 46]. The 60%SF₆-40%N₂ data are shown by the broken line.

gas-insulated equipment. However, more work is needed on particle-contaminated SF₆-N₂ gas mixtures. Present data are not sufficient to fully characterize the behavior of such mixtures as a function of partial or total pressure.

The solid-gas interface has its own breakdown characteristics. Generally, the breakdown voltage is lowered by the presence of a spacer. A 50%SF₆-50% N₂ mixture has about 90% the flashover voltage of pure SF₆ in the presence of spacers [31] (see also [53]). Similar conclusions have been reached by other researchers. For instance, Nakanishi [40] cited work by M. Hara et al. on the flashover characteristics of a cylindrical spacer on which a conducting particle was affixed in an SF₆-N₂ gas mixture which “showed that the flashover properties can be improved by mixing 20%SF₆ into N₂.”¹³

Furthermore, Eteiba et al. [54] measured the breakdown voltage of particle-contaminated spacers using 60-Hz and negative lightning impulse voltages. Figure 9 compares the dielectric strength of a clean spacer (curves A) to that obtained with a 2-mm long aluminum wire of 0.8 mm diameter particle attached on the spacer interface 2.6 mm from the center conductor (curves B) and from the outer envelope (curves D). The authors concluded that their results show no significant difference in the 60-Hz withstand voltage between pure SF₆ and the 50%SF₆-50%N₂ mixture for either the clean or the particle contaminated system and the 50%SF₆-50%N₂ mixture has only a slightly lower impulse ratio than pure SF₆. However, the breakdown characteristics of spacers for SF₆ and SF₆-N₂ mixtures as well as their cooling efficiencies need further investigation in order to quantify the performance of the mixtures under practical conditions, as has been pointed out by Endo [55].

¹³M. Hara et al. as quoted by Nakanishi [40].

TABLE 2. DC breakdown voltages of SF₆-N₂ mixtures with cylindrical electrodes and particle contamination [52]

SF ₆ -N ₂	V _{br} (kV) 303.9 kPa	V _{br} (kV) 405.2 kPa	V _{br} (kV) 607.8 kPa	V _{br} (kV) 810.4 kPa	V _{br} (kV) 1013 kPa
100/0	59.6	62.1	70.0	67.5	60.0
80/20	45.7	49.0	59.6	58.6	50.1
60/40	50.7	54.1	66.0	62.4	61.2
40/60	24.3	39.5	55.7	50.9	43.5
20/80	26.3	38.5	37.2	41.9	33.4

Similarly Blankenburg [56] found that the flashover behavior of cylindrical insulators in SF₆-N₂ mixtures subjected to AC voltage was found [88] to be qualitatively similar to that in pure SF₆ independently of the amount of SF₆ in the mixture (even when the mixture contained as little as 1% SF₆).

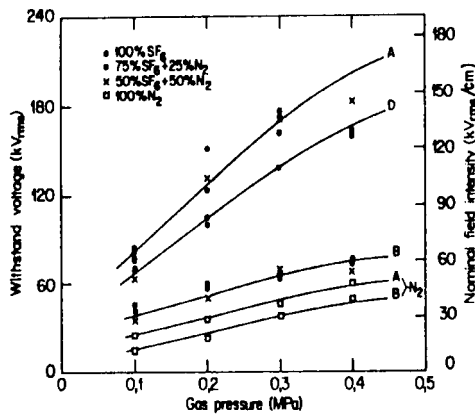
Additionally, according to Waymel and Boisseau [57] recent dielectric tests on real-size gas insulated transmission line (GITL) compartments confirmed the good industrial performance of SF₆-N₂ mixtures. Partial conclusions could be reached by them in the particular case of buried 400 kV GITL. They concluded that “compared to pure SF₆, SF₆-N₂ mixtures offer good compromise with the diameter of busbar required by the thermal design of buried GITL.”

Interestingly, the uniform field breakdown strength of SF₆ has been found [58] to increase by ~11% in the temperature range 300 K to 600 K. Because the presence of N₂ in SF₆-N₂ mixtures is not expected to change the dissociative electron attachment properties of SF₆ which are responsible for this increase, a similar increase in the dielectric strength of SF₆-N₂ mixtures with increasing temperature might be anticipated.

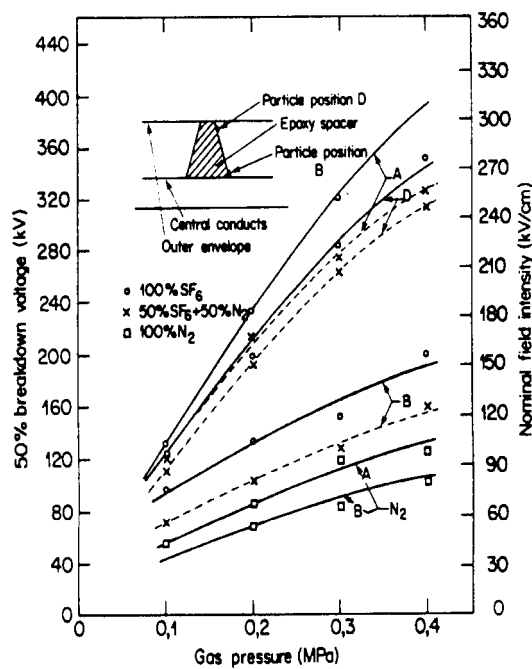
The behavior of a 50%SF₆-50%N₂ mixture under fast transient conditions has been investigated by Pfeiffer and co-workers. They found no significant difference in the behavior of the mixture compared to pure SF₆ (e. g, see Pfeiffer et al. [59] ; see also discussion in [6]).

Concerning the decomposition of SF₆-N₂ mixtures, there exist only limited data on the decomposition products of SF₆-N₂ mixtures (see discussion in [6]). These data are mainly of corona decomposition. They show that there is very little chemical interaction between SF₆ and N₂ in discharges and that the predominant oxidation byproducts are those seen in SF₆ (SO₂, SOF₂, SO₂F₂, and SOF₄). However, the presence of N₂ may reduce the ability of SF₆ to reform itself following an arc or a discharge and thus it may inhibit its recovery. This needs further investigation.

Finally, to our knowledge there is only one toxicological study [60] on SF₆-N₂ mixtures, which



(a)



(b)

FIG. 9. (a) Variation of the AC voltage in SF_6 , N_2 , and the mixtures 50% SF_6 -50% N_2 and 75% SF_6 -25% N_2 (from Eteiba et al. [54]). (b) Variation of the 50% negative impulse breakdown voltage for SF_6 , N_2 , and the mixture 50% SF_6 -50% N_2 (from Eteiba et al. [54]).

concentrates on the toxicological action of arc-decomposed SF_6 - N_2 mixtures (and pure SF_6). Its main conclusions are: (i) the lung, liver, and kidney can be attacked by arc-decomposed SF_6 gas and the alimentary system also can be influenced by the arced SF_6 ; and (ii) arced SF_6 - N_2 mainly attacks the lungs of the animals

exposed to the gas; livers, kidneys, and other organs are hardly affected, but the alimentary systems are generally influenced.

For gas-insulated transmission lines, SF_6 - N_2 mixtures in general and the 40% SF_6 -60% N_2 mixture in particular have great industrial potential. Depending upon the test, mixtures of SF_6 - N_2 appear to perform at least at levels of 80% of pure SF_6 , and in some cases SF_6 - N_2 mixtures exceed pure SF_6 performance. Indeed, some members of the electric power industry are in the process of designing and/or building GITL using SF_6 - N_2 mixtures for insulation purposes. The use of SF_6 - N_2 mixtures in existing GITL designed for pure SF_6 is more problematic due to the need for possible equipment changes and derating or recertification of existing equipment.

4.1.2 Gas-insulated Transformers

As was indicated in Sec. 2.4.2.2 of this report, heat dissipation is a significant requirement for gas-insulated transformers (GIT) in addition to the gas dielectric insulation characteristics. A number of recent studies [55, 61–65] considered SF_6 - N_2 mixtures as the insulating and heat transfer medium for gas-insulated transformers in spite of the fact that their insulating and heat-transfer (cooling) capabilities are somewhat lower than for pure SF_6 . A recent study on the temperature distribution in SF_6 - N_2 mixtures-insulated existing transformers [63] has led to the conclusion that “ SF_6 - N_2 mixtures-GIT meets the standards completely and it can be applied in electric power systems.” This same study found that a mixture ratio 55% SF_6 -45% N_2 has “good characteristics.” Similar studies [61, 62] concluded that: (i) the application of SF_6 - N_2 mixtures as the insulating and heat-transfer medium is feasible, (ii) with the same construction as for pure SF_6 , SF_6 - N_2 mixtures can be selected with composition as high as 55% SF_6 -45% N_2 with good insulation characteristics, and (iii) for 10 kV class SF_6 - N_2 mixtures GIT, the heat-transfer characteristics of the gas mixtures are the controlling factor in the design of insulation construction.

Because the temperature rise [55, 64, 65], of a 50% SF_6 -50% N_2 mixture over that of pure SF_6 is significant (approximately 15 °C to 20 °C) [63], it was suggested [64] that amorphous steel construction may help alleviate the heat transfer problem and allow use of a lower percentage of SF_6 in the SF_6 - N_2 mixtures. The use of SF_6 - N_2 mixtures may, thus, need to be coupled to the use of more heat resistant materials and modification of the transformer cooling design.

Overall, in spite of the difficulties mentioned in this section regarding the cooling capabilities of the SF_6 - N_2 mixtures, a 50% SF_6 -50% N_2 mixture can be a potentially useful gas-insulated transformer medium and further studies are indicated.

4.2 Interruption¹⁴

As discussed in Sec. 2, for arc and current interruption, the dielectric, switching, and thermal properties of the gas are important. Characterization of the cooling capacity of an interrupting gaseous medium involves consideration of the specific heat and specific thermal conductivity of the gas, as well as its ability to dissipate heat by convection. Nitrogen (and other light gases such as He) supplement SF₆ in this regard, SF₆ being efficient at the very high temperatures (say, 10,000 K to 3,000 K) and N₂ (or He) at the relatively lower temperatures (say, below 3,000 K). Table 3 lists the specific heats, specific thermal conductivities, and coefficients of viscosity of SF₆, N₂, and He.

There have been a number of studies on the arc and current-interruption capabilities of SF₆-N₂ mixtures and their performance in comparison to pure SF₆. Most such studies on circuit breakers used either of two types of gas circuit breakers (GCB). One is a double pressure type, and the other is a puffer type. Their structures and thus current-interruption capabilities are different. In the double pressure type GCB, high pressure gas is always stored in a high pressure vessel and the compressed gas in this vessel is blasted as soon as the contacts are separated. In the puffer type GCB, it is necessary to compress a gas in a puffer chamber during the opening stroke. The compressed gas is blasted to arc through an insulation nozzle. In this type of operation, the pressure increase and its duration are crucial variables for current-interruption capability. Most current GCBs are of the puffer type.

A comprehensive review and discussion of gases for arc interruption prior to 1982 was given in [3]. One of the significant results of this study is the observed strong dependence of the arc interruption performance of a gaseous medium on the total gas pressure, P . The rather limited data in this report [3] indicated that the arc interruption capability of SF₆ increased superlinearly with increasing pressure. This is significant because a small increase in the total pressure of an SF₆-N₂ mixture may compensate for the reduction in the arc interruption capability of the mixture relative to pure SF₆. This work [3] also concluded that: “For general purpose high voltage gas circuit breaker applications, SF₆ will be the interruption medium. However, there are applications for which other media can be viable alternatives. For example, a European manufacturer offers SF₆-N₂ as part of the puffer line with about 20% current derating in

TABLE 3. Specific heat, specific thermal conductivity, and coefficient of viscosity for SF₆, N₂, and He.

Gas	Specific heat ^a (cal g ⁻¹ K ⁻¹)	Thermal conductivity ^a (W m ⁻¹ K ⁻¹)	Coefficient of viscosity ^b (poise)
SF ₆	0.157	0.0155	161 x 10 ⁻⁶ (25 °C)
N ₂	0.248	0.0238	163 x 10 ⁻⁶ (0 °C)
He	1.242	0.150	189 x 10 ⁻⁶ (0 °C)

^aData provided by Endo [55] (Toshiba Corporation).

^bFrom Clark [71].

interrupting capability for applications in extreme low temperature environment (< -40 °C). Our data confirms that our puffer interrupters can be applied without design alteration. Furthermore, with interrupters designed specifically for SF₆-N₂, no derating is necessary with benefits of less SF₆ gas required, elimination of special heaters, and because of overall lower gas mass for a given pressure level, lower mechanical energy to operate the breaker. The combination can offer a lower cost interrupter with wider operating temperature range. Additional data on interrupter development specifically for SF₆-N₂ will be required to be more quantitative in regard to the economic advantages of SF₆-N₂ interruption medium.”

Another significant study is that of Grant et al. [72] who compared the performance of SF₆-N₂ mixtures as interruption media of gas-blasted arcs for various mixture compositions and total pressures of 500 kPa, 600 kPa, and 700 kPa. They measured the rate of rise of the recovery voltage (RRRV) capability, as a function of concentration of added N₂ (or He) to SF₆. Their results along with similar measurements by Garzon [73] are shown in Fig. 10. They show that the peak in the RRRV versus SF₆ percentage moves towards lower SF₆ concentrations at higher total pressures. These investigations also showed that the addition of appropriate amounts of N₂ (or He) to SF₆ can result in improved RRRV performance of up to 40% above that of pure SF₆ (Fig. 11). The measurements of Grant et al. [72] on SF₆-N₂ mixtures and Garzon [73] on SF₆-N₂ and SF₆-He mixtures are listed in Table 4. The measurements of Leeds et al. [74] on SF₆-air mixtures are also listed in Table 4 for comparison. As can be seen from Table 4, the measurements of Garzon [73] on the rate of rise of recovery of voltage (RRRV) for a synchronous interrupter show that the performance of SF₆-N₂ mixtures having 50%SF₆ by volume at pressures of 1300 kPa to 1900 kPa is approximately 1.39 times better than for pure SF₆. Garzon also found that the recovery capability of a non-synchronous circuit breaker using this gas mixture was at least as good as when pure SF₆ was used. The optimum interrupter performance, judged in terms of its voltage recovery capability, is observed to

¹⁴See Ref. [3] for a list of patents up to 1980 on gases for electrical arc interruption. Also see Chervy et al. ([66, 67, 68]) for information on the arc interruption capabilities of SF₆-CF₄ and SF₆-C₂F₆ mixtures, Nakagawa et al. [69] for the interruption capability of SF₆-CF₄ mixtures in puffer type gas-blast circuit breakers, and Middleton et al. [70] for work on SF₆-CF₄ circuit breakers.

TABLE 4. Recovery performance factors^a normalized to pure SF₆ (as listed in [69]).

%SF ₆	N ₂ 500 kPa [72]	N ₂ 600 kPa [72]	N ₂ 700 kPa [72]	N ₂ 1700 kPa [73]	He 600 kPa [73]	Air 1000 kPa [74]
100	1.0	1.0	1.0	1.0	1.0	1.0
90	1.32	1.07	1.13	1.02	0.98	-
75	1.03	1.24	1.46	1.08	1.12	-
65	-	-	1.52	1.17	-	-
60	-	0.93	-	1.22	1.13	-
50	1.0	0.82	0.86	1.33	1.14	0.52
40	-	-	-	1.39	-	-
25	0.56	0.38	0.52	0.90	1.08	0.28

^aRatio of the RRRV for a given mixture to the RRRV for pure SF₆.

occur when the mixture composition is roughly 50%SF₆-50%N₂ (Fig. 10). Garzon pointed out that these results using a double pressure type cannot be generalized for the design of all circuit breakers, but in applications where conditions are similar to those of his experiments, “it will be safe to assume that the use of a 50%-50% mixture of N₂ and SF₆ will serve to improve the dv/dt recovery capability of the interrupter.” While this conclusion is for higher pressures than are normally encountered in practice, Grant's data in Fig. 10 and the data in Table 4 clearly support this statement for lower total pressures as well.

The results of Garzon were obtained with a synchronously operated interrupter, and “therefore it is conceivable that non-synchronous operation may alter some of the findings.” However, Garzon states that “it is our experience that the recovery capability of a non-synchronous breaker using a 50%SF₆-50%N₂ mixture was at least as good as when 100%SF₆ had been used.” Their results taken between 1.38 MPa and 1.93 MPa indicate improved capability with increasing total pressure and this finding is consistent with earlier results [3].

Studies on full-size puffer interrupters using pure SF₆ and SF₆-N₂ mixtures by Sölver [75] led him to conclude that “a mixture of 69% SF₆ and 31% N₂ had considerably higher recovery speed than pure SF₆ at the same SF₆ partial pressure.”

As mentioned previously, Malik and Qureshi [38] reviewed the work on electrical breakdown in mixtures of SF₆ and other gases including N₂. They pointed out that previous work [72, 73, 76–78] shows that it is possible to further enhance the excellent arc interruption properties of SF₆ by using SF₆ mixed with lighter gases such as nitrogen or helium

Naganawa et al. [79] investigated the DC interruption by a spiral arc in SF₆-N₂ gas mixtures in the pressure range 0.1 MPa to 0.8 MPa. The mixtures they

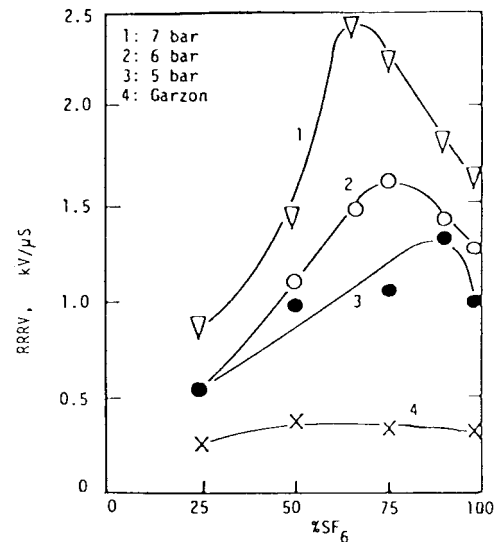


FIG. 10. RRRV as a function of SF₆-N₂ mixture ratio. Curves 1-3 are the data of Grant et al. [72] and Curve 4 are the data of Garzon [73] (adapted from Fig. 2 of [72]). The measurements of Garzon were made at a pressure of 1700 kPa.

studied contained a constant partial pressure of SF₆ equal to 0.1 MPa. The voltage-current characteristic curve of the spiral arc for the mixture 50%SF₆-50%N₂ was slightly below the similar curve for pure SF₆ at the same total pressure (0.8 MPa). They recommended SF₆-N₂ as “an extinguishing medium of switch gear to avoid the liquefying phenomena of pure SF₆ gas of high pressure under extremely low temperature and to reduce gas costs.”

However, other studies (see below) indicated that the 50%SF₆-50%N₂ mixtures performed not as well as pure SF₆ as arc or current interrupting media.

A comprehensive evaluation of and measurements on gases for arc interruption (puffer-type interrupter, current range ~10–15 kA) was conducted by Lee and Frost [80]. They concluded that “the results of previous investigators reaffirmed the overall excellent arc interruption ability of SF₆, while other gases and gas mixtures can have comparable performance in some aspects of interruption.” They themselves screened about 250 gases and out of these they selected 40 gases and gas mixtures for experimental evaluation. In Table 5 are given the arc interruption capabilities they measured for SF₆-N₂ mixtures, SF₆-He mixtures, and pure SF₆ for two values of the load line Z₀. These data show that the relative interruption capability of a 50%SF₆-50%N₂ mixture is only about 70% that of pure SF₆. This seems to be at variance with the studies mentioned above and points to the need for further studies.

Nakagawa et al. [69] performed calculations aimed at examining SF₆-N₂ gas mixtures in a buffer-type GCB. Their theoretical study showed that (i) the current interruption capability of the mixture depends on the

TABLE 5. Measured arc interruption capabilities^a of gases and gas mixtures at 0.6 MPa (Lee and Frost [80])

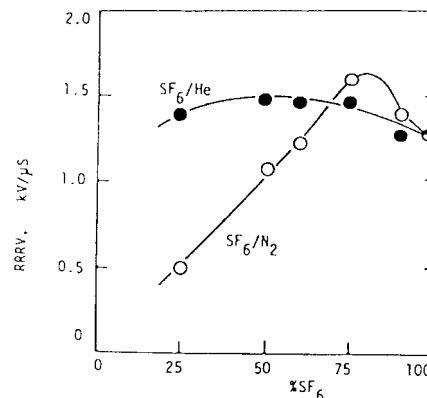
Gas or mixture	$Z_0 = 450 \Omega$ I_c (kA)	$Z_0 = 450 \Omega$ Relative interruption capability	$Z_0 = 225 \Omega$ Relative interruption capability	$Z_0 = 225 \Omega$ I_c (kA)
100% SF ₆	21.0	100	100	26.3
75%SF ₆ 25%N ₂	17.8	85	78	20.4
50%SF₆ 50%N₂	14.9	71	65	17.2
75%SF ₆ 25%He	15.4	73	78	20.4
50%SF ₆ 50%He	14.7	70	75	19.7

^aThe critical current I_c can be defined as that current where the critical RRRV line intersects the load line Z_0 . Higher I_c corresponds to higher interruption capability. For practical transmission and distribution power circuits, a Z_0 of 450 Ω is typical.

partial pressure of SF₆ in the mixture, and (ii) that SF₆-N₂ gas mixtures are inferior to pure SF₆. The interrupting ability of the SF₆-N₂ gas mixtures containing a given amount of SF₆ was found to deteriorate when N₂ was added to the mixture. For instance, the interrupting capability of 300 kPa of pure SF₆ was higher than that of a mixture of 300 kPa SF₆ + 200 kPa N₂. The no-load characteristics of 300 kPa SF₆ + 200 kPa N₂ gas mixtures showed that the rate of pressure rise was higher than that of pure 300 kPa SF₆, and that in spite of the higher value of the total pressure of the SF₆-N₂ mixtures, the pressure fall occurred faster in the mixture than in pure SF₆.

These findings are at variance with the work of Grant et al. [72] who reported that the interrupting abilities of SF₆-N₂ gas mixtures become higher at certain mixture ratio (see Figs. 10 and 11) but are consistent with the calculations of Tsukushi et al. [81] who examined the current interruption capability of SF₆ gas mixtures using puffer-type GCB. According to Tsukushi et al. for currents of ~15 kA, a 300 kPa SF₆ + 200 kPa N₂ showed 76% of di/dt of pure SF₆. Their calculation of the puffer pressure rise of gas mixtures in puffer-type gas blast circuit breakers for SF₆ and SF₆-N₂ mixtures indicated that the SF₆ partial pressure in a mixture was lower than the pure SF₆ pressure when the pure SF₆ filling pressure equaled that of the SF₆ partial pressure in the mixture. This was attributed to increases in the mass flow of SF₆ caused by the N₂ gas. Thus these calculations showed that the pressure characteristics in a puffer chamber are different for SF₆ and SF₆-N₂ mixtures. This seems to be born out by other calculations discussed below [82–87].

The interruption capability of SF₆-N₂ mixtures has

**FIG. 11.** RRRV as a function of SF₆-N₂ and SF₆-He mixture ratio for an upstream pressure of 600 kPa (from Grant et al. [72]).

been investigated by Gleizes et al. [82–87] in a series of papers. Specifically, Gleizes et al. [82–84] reported measurements of the axial temperature in a steady state arc plasma burning in SF₆-N₂ as a function of current intensity. They found that the axial temperature value is a complex function of radiation, thermal, and electrical conductivities and it may not be intermediate to those of pure SF₆ and pure N₂. At high currents the energy losses were found to be dominated by radiation. In another paper Gleizes et al. [81] calculated thermodynamic properties and transport coefficients for SF₆-N₂ mixtures in the temperature range 1000 K to 3000 K under the assumption that the number densities involved in the computation are those of a plasma under local thermodynamic equilibrium. Figure 12 shows some of their results on the dependence of the thermodynamic properties of N₂, SF₆, and SF₆-N₂ mixtures. It seems that the thermodynamic properties of the 40%SF₆-60%N₂ are not significantly different than those of pure SF₆.

Gleizes et al. [84] also performed calculations on the variations of temperature and conductance during the extinction of nonblown, atmospheric pressure, wall-stabilized arcs and concluded that “the use of SF₆-N₂ mixtures as a gas fill for circuit breakers will be efficient (i.e., will largely preserve the interruption properties of SF₆) when the proportion of SF₆ is higher than 50%.” See Refs. [85–87] for further calculations on the various parameters of significance in the performance of gas circuit breakers depending on type and gas medium and on the role of plasma convection.

Sasao et al. [88] simulated the arc dynamic behavior of gas-blasted arcs using SF₆-N₂ mixtures. Their simulations indicate that the use of SF₆-N₂ mixtures may require design changes of the arc chamber in order to optimize the arc quenching capability, and that these changes would depend on gas composition. They did not, however, indicate the “optimum” mixture composition. They found that the arc quenching ability of the SF₆-N₂

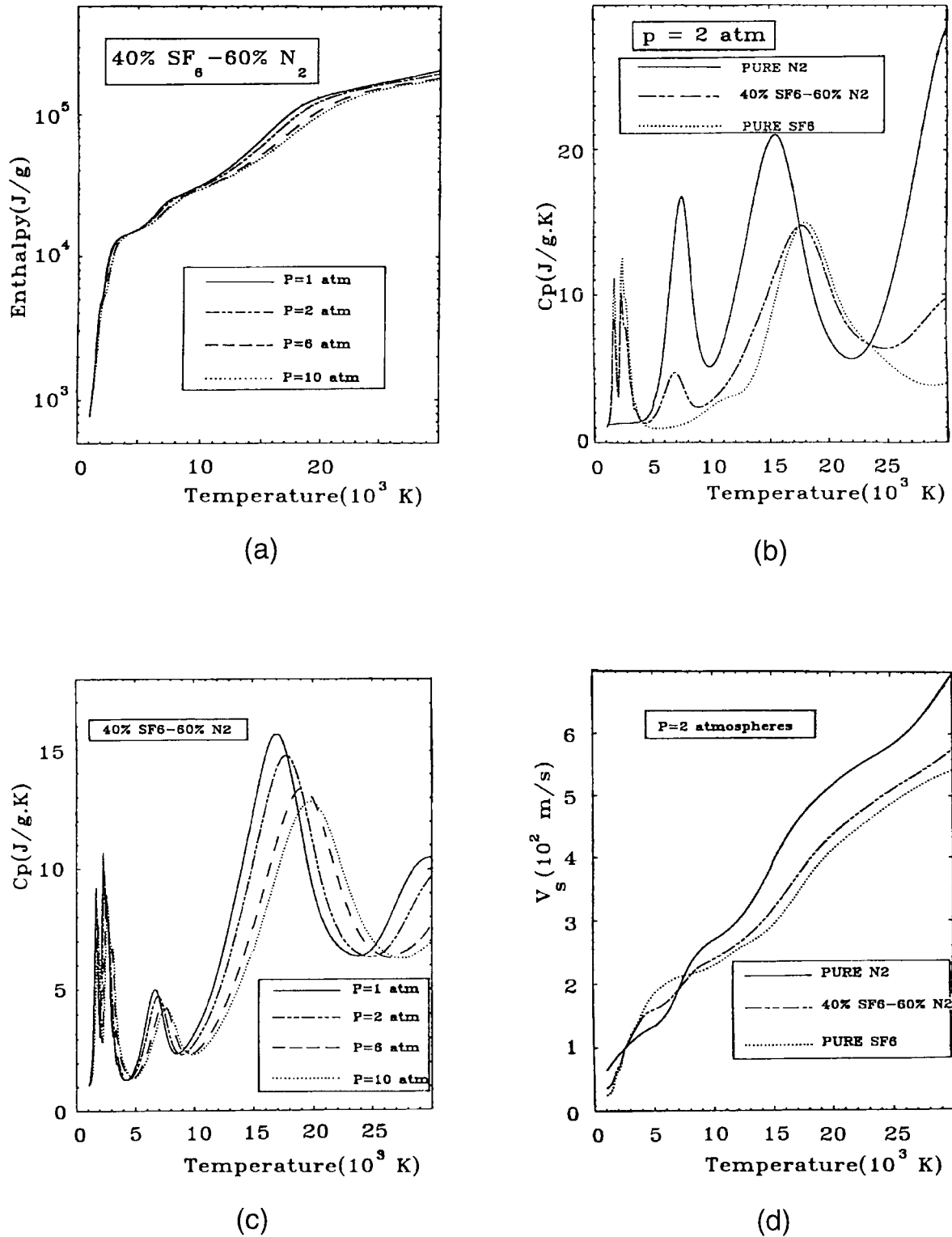


FIG. 12 A, B, C, D. Calculations by Gleizes et al. [83] of the variation of enthalpy with temperature and pressure of a 40%SF₆-60%N₂ mixture (Fig. 12a); variation of the constant-pressure specific heat, C_p, with temperature for pure N₂, pure SF₆, and 40%SF₆-60%N₂ mixture (Fig. 12b); evolution of the peaks of the constant-pressure specific heat with pressure of a 40%SF₆-60%N₂ mixture (Fig. 12c); and variations of the speed of sound, v, as a function of temperature for pure N₂, pure SF₆, and 40%SF₆-60%N₂ mixture (Fig. 12d). Note that these parameters for pure SF₆ and the 40%SF₆-60%N₂ mixture are rather close.

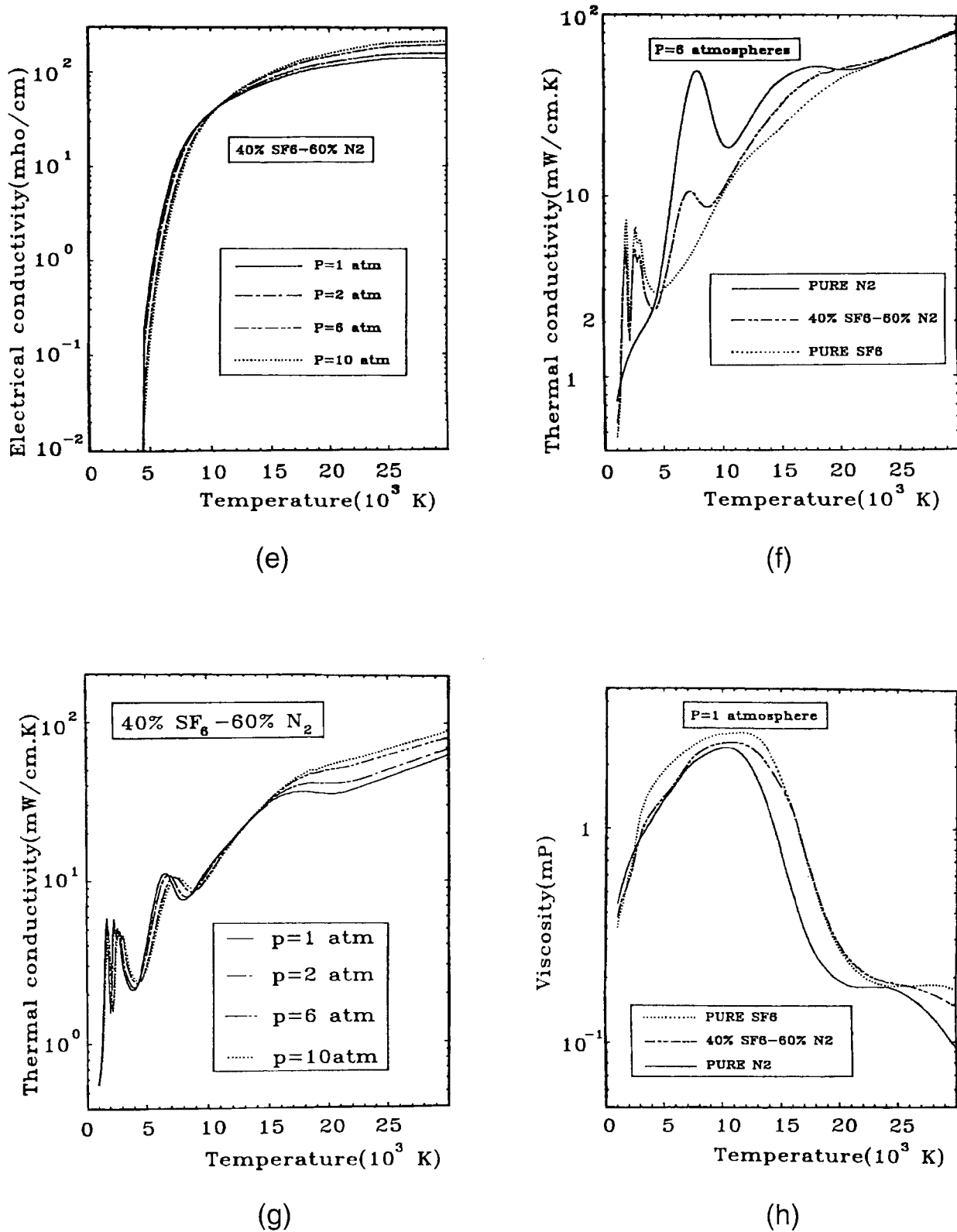


FIG. 12 E, F, G, H. Calculations by Gleizes et al. [83] for the electrical conductivity of a 40%SF₆-60%N₂ mixture as a function of temperature and pressure (Fig. 12e); variation of the total thermal conductivity with temperature for pure N₂, pure SF₆, and 40%SF₆-60%N₂ mixture at a total pressure of 6 atm (Fig. 12f); variation of the total thermal conductivity with temperature and pressure for 40%SF₆-60%N₂ mixture (Fig. 12g); and variation of viscosity with temperature for pure N₂, pure SF₆, and 40%SF₆-60%N₂ mixture at a total pressure of 1 atm (Fig. 12h).

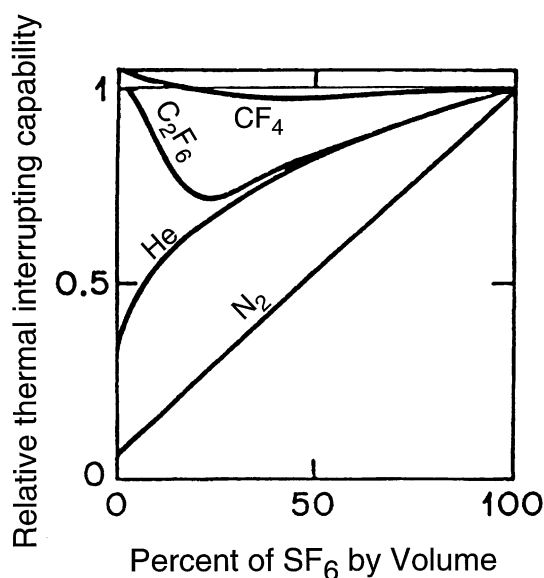


FIG. 13. Relative thermal interrupting capability of mixtures of He, N₂, CF₄, and C₂F₆ with SF₆ [70].

mixtures (including pressure rise and decrease) depends significantly on the configuration of the arc chamber and interruption current in addition to the arc quenching properties of the mixture itself.

Hence, these studies indicate that the actual performance is a function of many design variables. Thus a “drop-in” gas mixture (i.e., a gas mixture for use in existing equipment) does not appear to be feasible for GCBs designed specifically for use with pure SF₆. However, the concept of new circuit breakers designed for use with a mixture, rather than for pure SF₆, must be explored and tested before the possibility of a replacement mixture for circuit breakers is ruled out.

According to Waymel and Boisseau [57], gas-insulated substation circuit breakers require high arc-breaking properties that are not compatible with N₂ or low percentage SF₆-N₂ mixtures and for this reason SF₆-N₂ mixtures are not considered for switchgear and other gas-insulated equipment existing or re-designed.

Similarly, Middleton et al. [70] concluded that the use of SF₆-N₂ mixtures for circuit breakers involves significant derating of the circuit breakers under short line fault because of their reduced thermal capability compared to pure SF₆. These authors reported the relative thermal switching capabilities of various gas mixtures shown in Fig. 13 which indicate a poor performance for N₂ and a good performance for CF₄. As other studies have shown, the build up pressure is higher for the SF₆-N₂ mixture than for the SF₆-CF₄ and both are higher than for pure SF₆. It may thus be inferred from the studies mentioned above that the performance deficiencies of SF₆-N₂ mixtures in circuit breakers are principally due to thermal effects.

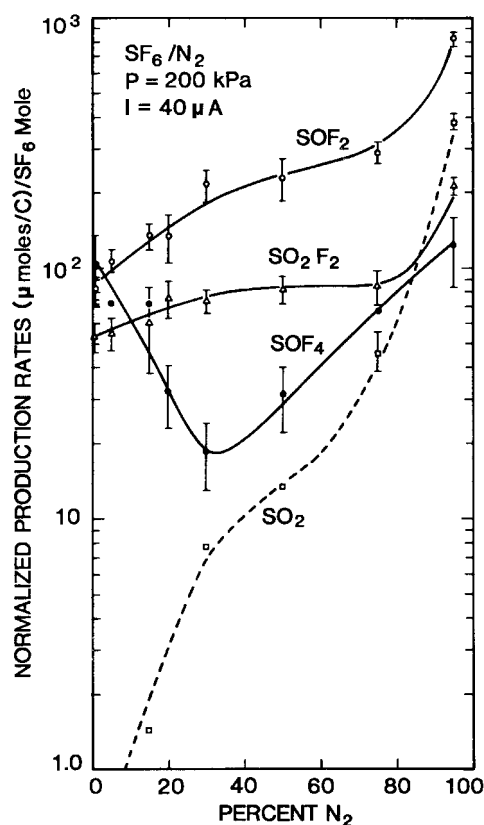


FIG. 14. Production rates normalized to the SF₆ content vs N₂ percent volume in SF₆-N₂ mixtures for negative point-plane corona in gas at a total absolute pressure of 200 kPa and a constant discharge current of 40 μA [6, 89].

Finally, Christophorou and Van Brunt [6] reviewed the limited data on the decomposition of SF₆-N₂ mixtures. Their conclusion, based on corona decomposition measurements, was that “there is little chemical interaction between SF₆ and N₂ in discharges, and the predominant oxidation byproducts are those seen in pure SF₆ such as SO₂, SOF₂, SO₂F₂, and SOF₄.” These byproducts are principally formed via interactions of SF₆ decomposition fragments with oxygen and water impurities [6]. The relative abundances of these byproducts may, however, be different, especially that of SO₂ which is much larger for the 40%SF₆-60%N₂ mixture compared to pure SF₆ (see Fig. 14). The very much larger concentrations of SO₂ in a 40%SF₆-60%N₂ mixture compared to pure SF₆ may be useful for diagnostic purposes. It might be noted also that the presence of N₂ may affect the ability of SF₆ to reform itself in arc or discharge decomposition.

Overall, the data for use of SF₆-N₂ mixtures in circuit breakers are contradictory, thus suggesting the need for additional research. It seems clear that SF₆-N₂ mixtures may not be used in existing breakers designed specifically for pure SF₆, but new designs may make effective use of SF₆-N₂ mixtures.

4.3 Gas Handling, Storing, Recycling, and Recovering

Clearly, if an SF₆-N₂ mixture is to be used in existing electrical equipment, a number of other issues need to be addressed and one of them pertains to information on handling, storing, recycling, and recovering SF₆ from the SF₆-N₂ mixtures. In this connection, Mitchel et al. [8] investigated the practical thermodynamics of SF₆ recovery from SF₆-N₂ (and SF₆-air) mixtures. They discussed recovery of SF₆ from mixtures with various non-condensable gases using a compressor/refrigerator system, and presented simulation results showing SF₆ recovery efficiency and capacity in terms of cooling temperature, total pressure, and gas composition. This study indicated that SF₆ extraction from a 50%SF₆-50%N₂ mixture presents no real problems. However, Probst [90] argues that SF₆-N₂ mixtures have problems in terms of recyclability and reusability and that economic factors may be significant (see Appendix C for additional discussion of this issue). The SF₆ gas can be reclaimed from the mixture but at a cost. Thus it seems reasonable to conclude that recycling of mixtures can be done, but the technologies used need to be improved. CIGRE 23.10 Task Force just published a document [19] dealing with SF₆ recycling, reuse of SF₆ gas in electrical equipment and final disposal. Unfortunately, no such work has been done on mixtures.

The data presented in earlier sections of this report suggest that there can be considerable tolerance for variation of the percentage of SF₆ in N₂ for a proposed 50%SF₆-50%N₂ mixture without significant effect on the dielectric performance of the mixture. This is because the properties of the mixture are not generally a strong function of the SF₆ concentration at this mixture composition. Certainly a tolerance in the percentage of SF₆ content of ±5% seems reasonable. It should also be noted that the removal of byproducts from the mixture is not expected to be much different than in pure SF₆. Furthermore, there seem to be no serious problems in making a standard gas mixture or in recovering the SF₆ from the mixture (see Appendix C).

4.4 Discussion

The electric power industry seems willing to consider SF₆-N₂ mixtures for insulation, for instance, in new gas-insulated transmission lines. Indeed, much work is being conducted world-wide in this area. Most such studies focus on low concentration mixtures (10% to 15%) for insulation, although work is also being done on higher concentration SF₆-N₂ mixtures for circuit breaker use. A 40%SF₆-60%N₂ mixture performs well as an insulating

medium. This “universal-application” mixture has no apparent physical or chemical problems, but the fact that its dielectric performance is only 85% to 90% of that of pure SF₆ would require equipment recertification along with hardware changes or derating. This is nearly impossible for equipment already in service, and would be costly for new equipment presently certified only for pure SF₆ use. Thus it would appear that the development of a replacement gas for use in existing equipment (a “drop-in” gas) is not presently a viable alternative. However, the application of standard gas mixtures to newly designed equipment is certainly worthy of further consideration.

Questions must also be raised with regard to the recovery, reusability, recycling, separation, and transportation of gas mixtures using existing technologies. These points are discussed in Appendix C of this report.

The electric power industry clearly prefers to use pure SF₆ for arc interruption. While still more work is necessary to resolve open questions and differences in published work, the standard mixture seems to have the potential to perform well even in circuit breakers, especially if used in new equipment designed specifically for use with a particular mixture. Nonetheless, it appears that industry is hesitant to consider SF₆-N₂ mixtures for arc interruption. Some of the reasons given [55, 91, 92], in addition to those mentioned above for insulation applications, are:

- Thermal derating would be required for many applications.
- The pressure rise during an internal failure arc in equipment will be much faster and higher with the mixture. This may be limited by rupture disk properties, which presents a possible safety issue.
- Some studies indicate significant reduction in the performance of mixtures, as compared to pure SF₆, in current circuit breaker designs, thus indicating the possible need for substantial breaker redesign.
- Recycling of mixtures will be more expensive and would require new equipment.
- Benefits of SF₆ substitutes can only be adequately judged by complete life cycle analysis of the equipment which is used, including the effects of different materials.

In general, the physical and chemical properties of a 40% or 50% mixture of SF₆ in N₂ suggest that it may be appropriate as a “universal application” gas mixture in new equipment, particularly if designed specifically for use with SF₆-N₂ mixtures. However, the practical difficulties of using SF₆-N₂ mixtures in existing equipment seem to be particularly large at present.

5. Other Promising Gases or Mixtures

In the previous section we have attempted to identify a gas mixture that would be acceptable as a “universal-application” replacement of SF₆ for both high voltage insulation and arc interruption. In this section we focus on gases or mixtures which are likely substitutes for specific high voltage insulation or arc interruption applications,¹⁵ and are thus worthy of immediate exploration (i.e., sufficient data are presently available to demonstrate their potential, but not to sufficiently prove their performance). Their possible use may require changes in equipment designs. We focus on three such gaseous dielectric media for which a significant amount of data are available:

- high-pressure pure N₂ for high voltage insulation;
- low concentration SF₆-N₂ mixtures for insulation and arc interruption; and
- SF₆-He mixtures for arc interruption.

Besides the gaseous media mentioned above, there are many other unitary gases, and binary and tertiary gas mixtures which are superior in dielectric strength to pure SF₆ and can potentially be used in high voltage needs (e.g., see [2, 3, 27, 31–33] and Table 1). However, the overwhelming preponderance of these gaseous dielectrics are not acceptable for various reasons such as their environmental impact, toxicity, or flammability, or because they cannot satisfy one or more of the required overall properties discussed in Sec. 2. From the long list of these we have identified a number of the most promising. These are discussed in Sec. 6.

5.1 High-Pressure N₂ for Insulation

As mentioned earlier [6, 14], nitrogen is an ideal gas to use: it is abundant, cheap, inert, non-toxic, non-flammable, and unquestionably environmentally acceptable.

However, nitrogen is a non-electronegative gas (it does not attach electrons) and for this reason its dielectric strength is rather low. Nitrogen, however, is a strong electron-slowng down gas and this property accounts for its relatively good dielectric properties in non-uniform fields and in the presence of conducting particles, and for its excellent performance in mixtures with electronegative

¹⁵Depending on a particular application, the mixture, composition, or pressure of the replacement gas will be varied to maximize the performance of the equipment.

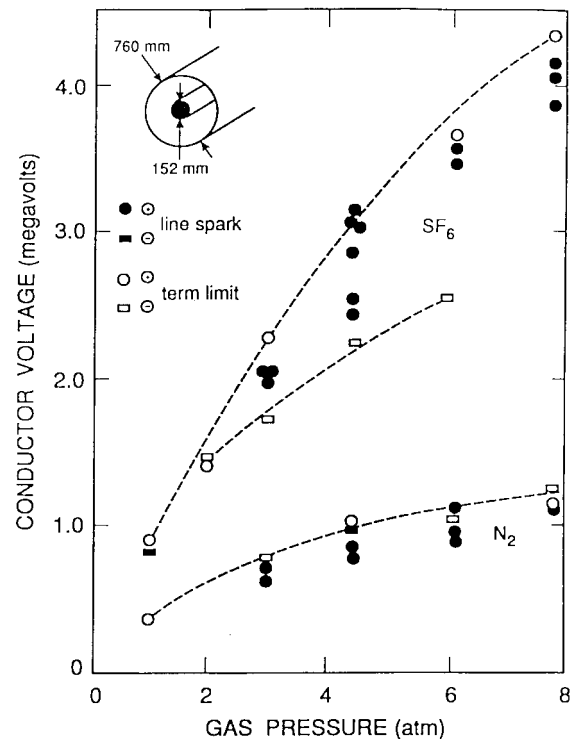


FIG. 15. DC breakdown voltage applied to the conductor as a function of gas pressure for SF₆ and N₂ using coaxial geometry (152 mm / 776 mm system; positive or negative polarity). The data represented by the solid and open symbols are for breakdowns at two locations as indicated in the figure. The solid symbols correspond to breakdown in the line and the open symbols are those cases where line sparks were not the limiting factor (see [95]).

gases [6, 25, 29]. Its thermal conductivity (Table 2) makes it a good cooling gas, especially at temperatures less than a few thousand degrees. In this regard, it nicely complements SF₆.

Existing measurements [3, 6, 14, 34, 37, 51] show that:

- Under uniform field conditions and low pressures (less than about 300 kPa) N₂ has about one third [25, 93] the dielectric strength of pure SF₆.

- The breakdown voltage (DC or AC) of N₂ increases with pressure as does that of SF₆ (see Figs. 4 and 15) [37, 48, 50, 94, 95], but it turns toward saturation at high pressures. The falling of the breakdown voltages for both N₂ and SF₆ below the linearly projected dielectric strength as the pressure increases, is due to the “magnification” at high pressures of the field non-uniformity due to surface roughness and imperfections. Such effects are more pronounced for SF₆ (and other electronegative gases) for which the effective ionization coefficient increases with the field much faster than does the ionization coefficient of the non-electronegative gas N₂ [6, 25, 93]. In Fig. 16 are shown the results of a recent comparison of AC and DC measurements using

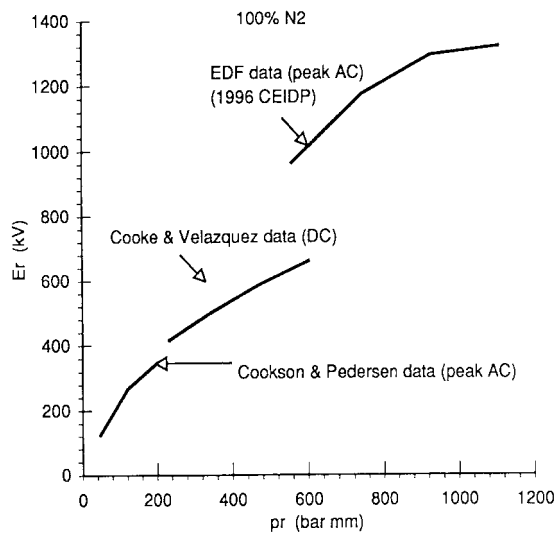


FIG. 16. Breakdown voltage E_r vs P_r (r is the radius of the inner conductor) for cylindrical electrode geometries (similarity plot) for pure nitrogen [34]. Data of Pace et al. [34], Cooke and Velazquez [95], and Cookson and Pedersen [37].

cylindrical electrodes.^{16,17,18} Pace et al. [34] argued that when the area effect is taken into consideration, the recent Electricité de France measurements [96] are compatible with the DC measurements of Ref. [37]. The limited lightning impulse measurements of Medeiros et al. [97] are consistent with the rest of the data in Fig.16. Breakdown voltages of 1 MV are possible for values of the product $P \times r$ (pressure times radius of inner conductor) of 8 MPa cm. The level of voltage is a function of the system dimensions and the total pressure. According to Pace et al., a rough estimate of the level of voltage may be determined by employing similarity rules.¹⁶

- Impulse breakdown studies [37, 96] with co-axial electrodes of various inner and outer conductor radii have been made and they vary with the ratio of the inner to outer conductor radius (see Fig. 17). The measurements of Cookson and Pedersen [37] with lightning impulse, are in excellent agreement with the recent measurements of

¹⁶We are thankful to M. O. Pace for Figs. 15, 16, and 19, and to X. Waymel and C. Boisseau for their permission to reproduce the EDF measurements.

¹⁷The similarity rule helps consolidate data from various experimental set ups. Two experiments are “similar” if one can be converted to the other by a change in scale. For example, two coaxial cylinder experiments are similar if the corresponding radii and lengths are all in the same ratio from one system to another [34, 98].

¹⁸The measurements of Refs. [36, 50, 94, 95] were made on coaxial geometries with various inner and outer conductor diameters.

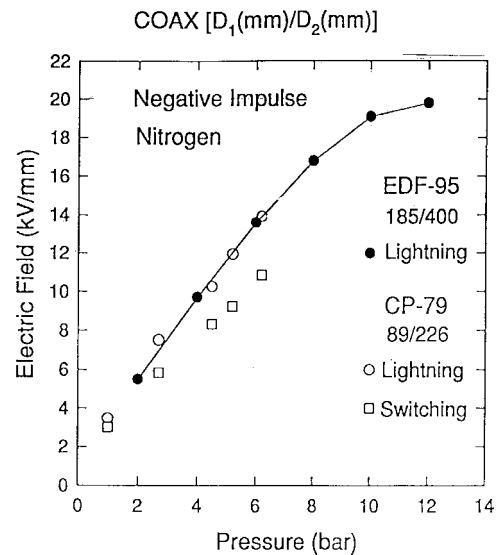


FIG. 17. Lightning impulse breakdown stress in nitrogen using cylindrical electrodes as a function of gas pressure [34, 96]. ●, Lightning data of Electricité de France [34, 96]; ○, Lightning data of Ref. 36; □, Switching data of Ref. 37. The ratio of the diameters of the two coaxial electrodes is given in the figure.

Electricité de France as reported by Pace et al. [34, 96]. For a coaxial electrode arrangement with 185 mm inner and 400 mm outer conductor radius, electric fields as high as 19 kV/mm can be sustained for nitrogen pressures of about 1MPa.

- The dielectric strength of N₂ is less sensitive to non-uniform fields than that of SF₆. This is understood from basic physical measurements such as the variation of the effective ionization coefficient with E/N close to $(E/N)_{lim}$ [14, 25, 93, 99, 100]. Similarly, N₂ is less sensitive than electronegative gases to conductor roughness. In practice, surface roughness effects are a strong function of the cable system size.

- Under conditions of conductive particle contamination and high pressures (about 1.0 MPa), N₂ performs very well compared to pure SF₆ (Fig. 8).

- The arc interruption capability of pure N₂ is significantly inferior to that of pure SF₆, although at high pressures (> 1 MPa) there may well be some use of pure N₂.

The physical data presented here suggest that high pressure (≥ 1 MPa) N₂ may be a good alternative to pure SF₆ for certain electrical insulation purposes. However, more work on practical systems at high pressures (high $P \times r$) is desirable to check its performance stability in industrial equipment. Also the question of environmental and economic impact of designing and constructing the required high pressure vessels must be investigated.

5.2 Low-Concentration SF₆-N₂ Mixtures for Insulation¹⁹

There have been many studies aimed at the development of nitrogen-based gaseous dielectrics. In Sec. 4 of this report we attempted to identify an “optimum” mixture of SF₆ and N₂ and in so doing we referenced many literature sources dealing with SF₆-N₂ mixtures as a function of SF₆ concentration. In this section we focus on the possibility of developing *low-concentration* SF₆-N₂ mixtures for possible use in electrical insulation. By low concentration is meant a percentage of SF₆ in N₂ of less than 15%. The available information on such mixtures is outlined below (see also, Sect. 4).

- Small amounts of electron attaching gases such as SF₆ in N₂ substantially increase the dielectric strength of the mixture (Fig. 4). Depending on the electron attaching properties of the electronegative gas which is added to N₂, the increase in the dielectric strength of the mixture may or may not saturate as the electronegative gas concentration is increased [25, 42].

- Pace et al. [34, 96] compared the AC measurements of Cookson and Pedersen [37] with the measurements of EDF for co-axial cables. Their results are shown in Fig. 18 for 5% and 10% mixtures. A similar comparison was made by them for the negative lighting impulse breakdown voltage as a function of pressure for a number of gas mixture compositions. An example of these measurements and comparisons are given in Fig. 19 for a 10% mixture. The data [34, 96] are in reasonable agreement when the similarity law for cylinders is applied (Fig. 19). It should be noted, however, that the increase in the breakdown voltage with pressure is not linear and any simple extrapolation to higher pressures of these results may be in error.

- Malik et al. [101] measured the breakdown properties of low concentrations (< 1.5%) of SF₆ in N₂ in a highly non-uniform field arrangement (rod-plane geometry). Their results for negative polarity clearly show a large increase in dielectric strength even at very low concentrations (< 0.3%). Such mixtures may be useful for filling substations. In such situations there are parts that have special requirements in terms of higher

¹⁹See, also, Volume 2 of the Proceedings of the 10th International Symposium on High Voltage Engineering, August 25–29, 1997, Montréal, Québec, Canada. For instance, H. I. Marsden, S. J. Dale, M. D. Hopkins, and C. R. Eck III, “High Voltage Performance of a Gas Insulated Cable with N₂ and N₂-SF₆ Mixtures,” pp. 9–12; T. B. Diarra, A. Bérroual, F. Buret, E. Thuries, M. Guillen, and Ph. Roussel, “N₂-SF₆ Mixtures for High Voltage Gas Insulated Lines,” pp. 105–108; and X. Waymel, V. Delmon, T. Reess, A. Gibert, and P. Domens, “Impulse Breakdown in Point-Plane Gaps in SF₆-N₂ Mixtures,” pp. 289–292.

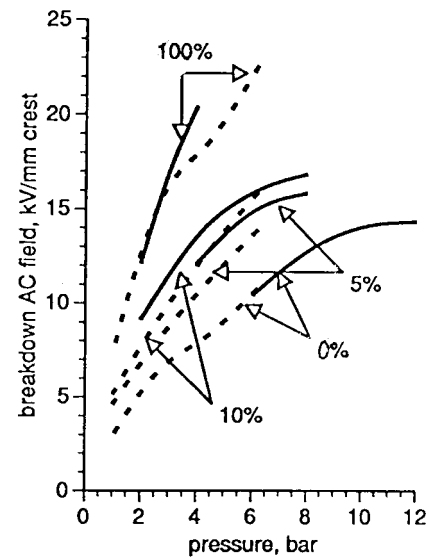


FIG. 18. Measured breakdown fields in coaxial cables of diameters 185 mm / 400 mm (EDF-95 [96], solid curves), and 89 mm / 226 mm ([37], dashed curves). The percentage of SF₆ in mixtures with nitrogen is indicated in the figure [34, 96].

levels of insulation. These can be separately insulated with gases or mixtures containing higher percentages of SF₆ or even with pure SF₆ if indeed this is necessary. They certainly can be used for transmission of lower level voltages.

- A lightning impulse (1.2 / 50 μs) study [102] of SF₆-N₂ mixtures with 0.15% to 0.2% SF₆ content for rod/plane gaps with both positive and negative voltages showed that for both polarities the effect of the addition of SF₆ to N₂ is dependent on both the gas pressure and gap spacing. Maxima in voltage versus SF₆ percentage curves were observed which were a function of the total pressure.

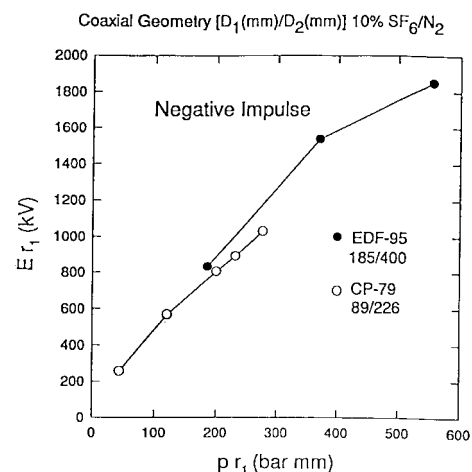


FIG. 19. The product E_r as a function of the product P_r for a 10% SF₆-90% N₂ mixture for lightning impulse breakdown (E is the electric field, r is the radius of the inner cylinder of the coaxial cylinder electrode geometry, and P is the total pressure) [34, 96].

For positive polarity voltages the maximum in breakdown strength occurs when the SF₆ content in the mixture is about 0.5% at 100 kPa, 5% at 300 kPa, and 10% at 500 kPa.

- Qiu and Kuffel [103] investigated the increase in dielectric strength of nitrogen (and helium) mixtures due to 1% SF₆ additive. Table 6 shows their data on the $V_{\text{mixture}} / V_{\text{gas}}$ of the breakdown voltage for the mixture to the breakdown voltage of the buffer gas (N₂, or He). It is seen that even 1% SF₆ in N₂ significantly improves the dielectric strength, and that this improvement varies with the type of applied voltage.

- Yializis et al. [104] studied impulse breakdown and corona characteristics of SF₆-N₂ mixtures with less than 1% of SF₆ content using rod-plane gaps. Measurements of 50% impulse breakdown voltage were made mainly in SF₆-N₂ mixtures containing 0.1% SF₆ content by pressure over the range of 100 kPa to 500 kPa and gap lengths of 10 mm to 50 mm using positive and negative polarity 1.8 / 50 ms and 310 / 3500 ms pulses. Their results show that the positive impulse breakdown of N₂ in the pressure region of 100 kPa to 250 kPa increases considerably with the addition of small traces of SF₆.

- Naganawa et al. [73] made measurements on DC interruption by spiral arc in SF₆-N₂ mixtures (0.1 MPa to 0.8 MPa). They recommended SF₆-N₂ mixtures as an extinguishing medium of switchgear to avoid liquefaction of pure SF₆ at high pressure under extremely low temperatures and to save on gas cost. They concluded that compared to the case of pure SF₆ “even a small content of SF₆ in the mixture is effective to decrease the magnitude of interrupting overvoltages with the arcing time unchanged.” On the other hand, Wootton and Cookson [51] found that “addition of trace amounts of SF₆ (e.g., 2%) to nitrogen at high pressures (e.g., 1.2 MPa) can reduce the breakdown strength (by ~40%), while increasing the strength at low pressures.

- Finally, according to Bolin [105], recent reports from ABB and Siemens show that GITL are being designed for use with low percentage SF₆ mixtures (containing less than 20% SF₆).

TABLE 6. $V_{\text{mixture}} / V_{\text{gas}}$ for a mixture of 1% SF₆ in either N₂ or He for various types of applied voltage [103].

Applied Voltage	$V_{\text{mixture}} / V_{\text{gas}}$ (400 kPa)	$V_{\text{mixture}} / V_{\text{gas}}$ (200 kPa)
	He	N ₂
-1.5 / 40 μs	1.35	1.55
+ 1.5 / 40 μs	1.92	1.70
AC	2.59	2.79

TABLE 7. Breakdown strength of mixtures of SF₆ and He [106].

Gas mixture	Strength relative to SF ₆ (quasi-uniform field ; sphere-plane electrodes at 150 kPa)	Maximum strength relative to SF ₆ (non-uniform field ; pin protrusion; 100 kPa < P < 500 kPa)	Minimum strength relative to SF ₆ (non-uniform field ; pin protrusion; 100 kPa < P < 500 kPa)
100% SF ₆	100	100	100
100% He	~3	-	-
75% SF ₆ 25% He	78	130	71
50% SF ₆ 50% He	56	126	58
25% SF ₆ 75% He	33	101	41

5.3 SF₆-He Mixtures for Arc Interruption

SF₆-He mixtures are considered for use in circuit breakers. Helium has a very low dielectric strength (~ 3% that of SF₆ in uniform fields, Table 7) and contributes virtually nothing to the dielectric strength of the mixture. Unlike the SF₆-N₂ mixtures which exhibit substantial synergism in terms of their dielectric strength, the SF₆-He mixtures show no such synergism.

Helium, however, complements SF₆ in terms of its cooling capability because it is very light. Its specific heat and thermal conductivity are very large (see Table 3). Helium is an inert gas and does not react chemically with either SF₆, or the gas impurities present in commercial SF₆, or the system components.

Grant et al. [73] investigated the recovery performance of SF₆-He mixtures as a function of the SF₆ content in the mixture. Their results were presented earlier in Table 4 and show the performance of SF₆-He mixtures which for a total pressure of 0.6 MPa seem to be ~10% higher than pure SF₆ for virtually all mixture compositions from 75% to 25% SF₆ (see Fig. 11).

Basile et al. [107] found that SF₆-He mixtures show a synergistic maximum in the 50% breakdown voltage at a percentage of He around 30%. At the maximum content of He used in these tests (50%), the breakdown voltage of a rod-plane electrode system was still higher than in pure SF₆. This suggests that SF₆-He mixtures could be considered as an alternative to pure SF₆ in operating conditions of low temperature.

Wootton and Cookson [51] measured the 60-Hz dielectric strength of SF₆-He mixtures in a plane-parallel electrode system with particle contamination (free 6.4 mm long x 0.45 mm diameter copper and aluminum particles).

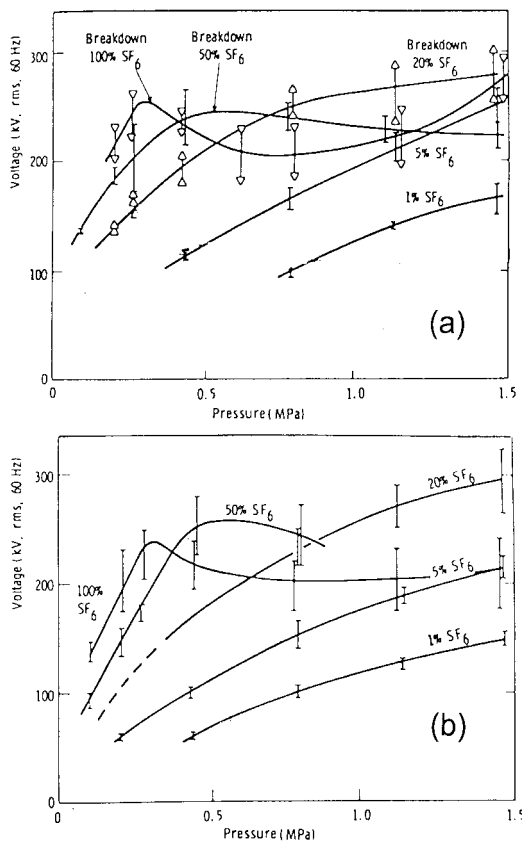


FIG. 20. Dielectric strength for SF₆-He mixtures with 1%, 5%, 20% and 50% SF₆ for copper (Fig. 20 a) and aluminum (Fig. 20 b) particles [51].

Figure 20 shows the dielectric strength for mixtures with 1%, 5%, 20% and 50% SF₆ for copper (Fig. 20a) and aluminum (Fig. 20b) particles. While the breakdown strength of helium in uniform fields is only a small fraction of that of SF₆ and mixtures of the two gases in uniform fields have dielectric strengths which are intermediate to those of the two gases, the data in Fig. 20 show that the dielectric strength of the He-SF₆ mixtures under particle contamination is equal to or greater than that of SF₆ at its optimum pressure of about 0.29 MPa. Further analysis of these data by Wootton and Cookson showed that the maxima in the curves for total pressures of 1.43 MPa, 1.1 MPa, 0.77 MPa, and 0.43 MPa occurred at a partial pressure of SF₆ of about 0.3 MPa. It was also found that for mixtures with the optimum partial pressure of SF₆ (0.29 MPa), addition of He increases the breakdown strength linearly, in contrast to addition of SF₆ which decreases the breakdown strength. The data in Fig. 20b show that a 20% SF₆-80% He mixture at total pressures greater than 0.7 MPa is superior in performance to pure SF₆. Even a 5% SF₆-95% He mixture at a total pressure greater than 1.3 MPa performs better than pure SF₆.

This work on particles – and the work discussed in

this and earlier sections (e. g., Table 4, Figs. 10 and 11) on arc interruption – would suggest that SF₆-He mixtures containing 0.29 MPa of SF₆ at total pressures in excess of about 1 MPa may be suitable for arc/current interruption applications. However, Niemeyer [108] points out that the published data on SF₆-He synergisms in switchgear only refer to the initial thermal recovery under short line fault switching where the dielectric stress is still low and not to terminal fault interruption where the dielectric stress is high. Clearly, there are still a number of issues regarding the SF₆-He mixtures which include gas compression, separation and leakage which need to be addressed.

The data presented here strongly suggest that pure N₂ and mixtures of low concentration of SF₆ in N₂ may be appropriate for many insulating applications. The data also suggest that SF₆-He mixtures may be viable for use in gas-insulated circuit breakers. Further research on their use in practical systems must still be performed.

6. Other Possible Substitutes: Future Long-term R&D

As mentioned previously, other gases exist for which there are data that indicate some potential for use as a gaseous dielectric or interruption medium. Some of those with the most potential are listed in Table 8, and an investigation of these gases could form the core of a future long-term research and development effort to develop acceptable substitute gaseous media for the various needs of the electric power industry. This program could include efforts in the following specific areas:

- A search for single gases at high pressure (besides N₂) for insulation purposes, including CO₂, N₂O, and SO₂ [2, 3, 6, 14, 30, 31, 109–111]. These are weakly electronegative gases and their electron attaching properties and dielectric strength may increase with gas density [110, 111].
- A search for binary mixtures (other than SF₆-N₂ and SF₆-He) suitable for particular applications [3, 2, 25, 30, 31, 42, 51, 53, 80, 112–123], including: SF₆ + Ar, SF₆ + CF₄, SF₆ + C₂F₆, and He + an electronegative component gas (for arc and current interruption); N₂ + SO₂, N₂ + c-C₄F₈, SO₂ + SF₆, SF₆ + CO₂, and N₂O + SF₆ (for insulation).

TABLE 8. Gases and gas mixtures, for possible use in insulation and arc and current interruption applications, which may be appropriate for further research based upon presently known information.

	Insulation	Interruption
Possible Universal Mixtures	<ul style="list-style-type: none"> • 40% SF₆ + 60% N₂ • 50% SF₆ + 50% N₂ 	<ul style="list-style-type: none"> • 40% SF₆ + 60% N₂ • 50% SF₆ + 50% N₂
Near Term Research	<ul style="list-style-type: none"> • High pressure N₂ • Low concentration SF₆ in N₂ 	<ul style="list-style-type: none"> • SF₆ + He
Long Range Research	<ul style="list-style-type: none"> • CO₂ • SO₂ • N₂O • N₂ + SO₂ • N₂ + c-C₄F₈^a • SO₂ + SF₆ • SO₂ + c-C₄F₈ • SF₆ + CO₂ 	<ul style="list-style-type: none"> • SF₆ + Ar • SF₆ + CF₄^a • SF₆ + C₂F₆^a • SF₆ + N₂ + He • SF₆ + N₂ + Ar • He + electronegative gases

^aThese are also greenhouse gases, but their global warming potentials are about one third that of SF₆. Over a 100-year time horizon the global warming potentials of c-C₄F₈, CF₄, C₂F₆, and SF₆ are respectively 8,700, 6,500, 9,200, and 23,900 [22].

- A search for ternary mixtures, including the following systems: SF₆ + N₂ + He and SF₆ + N₂ + Ar [3, 2, 30, 31, 124, 125].

While a large amount of effort was expended in the 1970s and 1980s to search for gases exhibiting better dielectric performance than SF₆, the emphasis of any new research program would be to identify gases with acceptable dielectric properties, and minimal environmental impact. This could include a search for new synthetic gases, better additives than SF₆, better buffers than N₂, and gases with an IR window near 10 μm to avoid any contribution to global warming. It could even be suggested that an investigation of gaseous media that are detrimental to the environment is justified, if the gas has superior performance properties for a particular application where its release into the environment could be absolutely prevented.

Other areas of productive investigation that are suggested by the research presented in this report include the following:

- the role of humidity and impurities on dielectric gas properties;
- electron attachment and detachment in low concentration SF₆-N₂ mixtures;
- decomposition of SF₆-N₂ mixtures as a function of concentration, impurity content, and type of discharge;
- dielectric behavior of gas mixtures at high gas pressures;
- thermal and electrical conductivities of SF₆-N₂ mixtures; thermodynamic properties, thermal

conductivities, and viscosities of gas mixtures under the conditions they are used in electrical equipment;

- the role of particles and improved particle control methods;
- interface phenomena and partial discharge behavior;
- assessment of available data;
- life cycle analysis of the overall environmental gain by using mixtures instead of pure SF₆ ;
- handling, storage, recovery, and disposal of gas mixtures;
- ways to reduce the cost of recovery of SF₆ from gas mixtures;
- new equipment designed specifically to eliminate emissions;
- improved recycling procedures;
- relaxation of equipment constraints that would make other gases acceptable;
- alternative technologies not requiring gaseous dielectrics, such as high temperature superconductors and solid state switching.

A significant amount of research must be performed for any new gas or gas mixture to be used in electrical equipment. Such a program necessarily would require the systematic study of potential replacements, including their physical, chemical, and performance properties. A concerted national or international effort in this area by equipment manufacturers, utilities, government labs, universities and gas manufacturing companies would be beneficial.

7. Conclusions and Recommendations

Sulfur hexafluoride is a superior dielectric gas for nearly all high voltage applications. It is easy to use, exhibits exceptional insulation and arc-interruption properties, and has proven its performance by many years of use and investigation. It is clearly superior in performance to the air and oil insulated equipment which was used prior to the development of SF₆-insulated equipment. However, the extremely high global warming potential of SF₆ mandates that users actively pursue means to minimize releases into the environment, one of which is the use of other gases or gas mixtures in place of SF₆.

An evaluation of the results of the last two decades, and a detailed analysis of the data presented in this report, indicate that no replacement gas is immediately available for use as an SF₆-substitute (“drop-in gas”) in existing electric utility equipment. For gas insulated transmission lines and gas insulated transformers, the limitation is primarily due to the need for re-certification and possible re-rating of equipment that is already in use. For gas insulated circuit breakers there are still significant questions concerning the performance of gases other than pure SF₆.

However, various gas mixtures show considerable promise for use in new equipment, particularly if the equipment is designed specifically for use with a gas mixture:

- Mixtures of nearly equal amounts of SF₆ and N₂ exhibit dielectric properties that suggest that they could be used as a “universal application” gas for both electrical insulation and arc/current interruption purposes. In this connection, standard procedures for mixture handling, use, and recovery would need to be further developed.

- Mixtures of low concentrations (<15%) of SF₆ in N₂ show excellent potential for use in gas insulated transmission lines, although further work on their performance in practical systems is necessary.

- Pure high pressure nitrogen may be suitable for some electrical insulation applications. Consideration of the use of such environmentally friendly gases where SF₆ is not absolutely required should be investigated and promoted.

- A mixture of SF₆ and helium has shown promise when used in gas insulated circuit breakers, and should be investigated further.

Finally, it is clear that a significant amount of research must be performed for any new gas or gas mixture to be used in electrical equipment. Such a program necessarily would require the systematic study of potential replacements, including their physical, chemical, and performance properties (see for example Sections 3 and 6). A concerted effort in this area by equipment manufacturers, utilities, government labs, universities, and gas manufacturing companies would be beneficial.

8. References

1. G. Camilli, “Gas-Insulated Power Transformers,” *Proc. IEE*, Vol. 107 A, pp. 375–382, 1960.
2. L. G. Christophorou (Ed.), *Gaseous Dielectrics*, Volumes I-V, Pergamon Press, New York, respectively, 1978, 1980, 1982, 1984, 1987; Volumes VI and VII, Plenum Press, New York, respectively, 1990 and 1994.
3. Electric Power Research Institute, “Gases Superior to SF₆ for Insulation and Interruption,” Report EPRI EL-2620, September, 1982.
4. S. A. Boggs, F. Y. Chu, and N. Fujimoto (Eds.), *Gas-Insulated Substations*, Pergamon Press, New York, 1986.
5. W. T. Shugg, *Handbook of Electrical and Electronic Insulating Materials*, Second Edition, IEEE Press, New York, pp. 433–452, 1995.
6. L. G. Christophorou and R. J. Van Brunt, “SF₆/N₂ Mixtures, Basic and HV Insulation Properties,” *IEEE Trans. Dielectrics and Electrical Insulation*, Vol. 2, pp. 952–1003, 1995.
7. A. H. Cookson, “Gas-Insulated Cables,” *IEEE Trans. Electr. Insul.* Vol. EI-20, pp. 859–890, 1985.
8. G. R. Mitchel, J. Castonguay, and N. G. Trinh, “Practical Thermodynamics of SF₆ Recovery from SF₆/N₂ or SF₆/Air Mixtures,” in *Gas-Insulated Substations*, S. A. Boggs, F. Y. Chu, and N. Fujimoto (Eds.), Pergamon Press, New York, pp. 437–442, 1986.
9. G. Mauthe, L. Niemeyer, B. M. Pryor, R. Probst, H. Bräutigam, P. A. O’Connell, K. Pettersson, H. D. Morrison, J. Poblitzki, and D. Koenig, Task Force 01 of Working Group 23.10, “SF₆ and the Global Atmosphere,” *Electra* No. 164, pp. 121–131, February, 1996.
10. A. Diessner, CAPIEL draft paper on “The Influence of Switchgear on the Greenhouse Effect,” Private Communication, 1996.
11. G. Mauthe, K. Pettersson, P. Probst, J. Poblitzki, D. Köning, L. Niemeyer, and B. M. Pryor, Members of WG23.10 Task Force 01. Draft document on “SF₆ and the Global Atmosphere,” March 1995.
12. “Sales of Sulfur Hexafluoride (SF₆) by End-Use Applications,” Science & Policy Services, Inc., The West Tower, Suite 400, 1333 H Street NW, Washington, D.C. 20005, 1997.
13. E. Cook, “Lifetime Commitments: Why Climate Policy-Makers Can’t Afford to Overlook Fully Fluorinated Compounds,” World Resources Institute, Washington, D.C., February, 1995.
14. L. G. Christophorou and R. J. Van Brunt, “SF₆ Insulation: Possible Greenhouse Problems and Solutions,” NISTIR 5685, July, 1995.
15. M. Maiss, and I. Levin, “Global Increase of SF₆ Observed in the Atmosphere,” *Geophys. Res. Lett.*, Vol. 21, pp. 569–572, 1994.
16. C. P. Rinsland, M. R. Gunson, M. C. Abrams, L. L. Lowes, R. Zander, and E. Mahieu, “ATMOS/ATLAS 1 Measurements of Sulfur Hexafluoride (SF₆) in the Lower Stratosphere and Upper Troposphere,” *J. Geophys. Res.*, Vol. 98, pp. 20 491–20 494, 1993.; V. Ramanathan, L. Callis, R. Cess, J. Hansen, I. Isaksen, W. Kuhn, A. Lacis, F. Luther, J. Mahlman, R. Reck, and M. Schlesinger, “Climate Chemical Interactions

- and Effects of Changing Atmospheric Trace Gases,” *Reviews Geophysics*, Vol. 25, pp. 1441–1482, 1987.
17. (a) P. K. Patra, S. Lal, B. H. Subbaraya, C. H. Jackman, and P. Rajaratnam, “Observed Vertical Profile of Sulphur Hexafluoride (SF₆) and Its Atmospheric Implications,” *J. Geophys. Res.*, Vol. 100, pp. 8855–8859, 1997; (b) L. S. Geller, J. W. Elkins, J. M. Lobert, A. D. Clarke, D. F. Hurst, J. H. Butler, and R. C. Myers, “Tropospheric SF₆: Observed Latitudinal Distribution and Trends, Derived Emissions and Interhemispheric Exchange Time,” *Geophys. Res. Lett.*, Vol. 24, pp. 675–678, 1997.
 18. M. K. W. Ko, N. D. Sze, W.-C. Wang, G. Shia, A. Goldman, F. J. Murcray, D. G. Murcray, and C. P. Rinsland, “Atmospheric Sulfur Hexafluoride: Sources, Sinks, and Greenhouse Warming,” *J. Geophys. Res.*, Vol. 98, pp. 10499–10507, 1993.
 19. G. Mauthe, B. M. Pryor, L. Niemeyer, R. Probst, J. Poblitzki, H. D. Morrison, P. Bolin, P. O’Connell, and J. Henriot, “SF₆ Recycling Guide”, CIGRÉ 23.10 TASK FORCE 01, April 1997.
 20. U. S. Environmental Protection Agency, “Electrical Transmission and Distribution Systems, Sulfur Hexafluoride, and Atmospheric Effects of Greenhouse Gas Emissions Conference,” EPA Conference Proceedings Report, August, 1995.
 21. D. Edelson and K. B. McAfee, “Notes on the Infrared Spectrum of SF₆,” *J. Chem. Phys.*, Vol. 19, pp. 1311–1312, 1951.
 22. Intergovernmental Panel on Climate Change (IPCC), “Climate Change 1995,” J. T. Houghton, L. G. Meira Filho, B. A. Callander, N. Harris, A. Kattenberg, and K. Maskell (Eds.), Cambridge University Press, Cambridge, New York, p. 22, 1996.
 23. R. A. Morris, T. A. Miller, A. A. Viggiano, J. F. Paulson, S. Solomon, and G. Reid, “Effect of Electron and Ion Reactions on Atmospheric Lifetimes of Fully Fluorinated Compounds,” *J. Geophysics Research*, Vol. 100, pp. 1287–1294, 1995.
 24. D. J. Wuebbles and A. K. Jain, “Concerns About Climate Change and SF₆,” In Ref. [20], pp. 8–38.
 25. L. G. Christophorou, “Insulating Gases,” *Nuclear Instruments and Methods in Physics Research*, Vol. A268, pp. 424–433, 1988.
 26. L. G. Christophorou and P. G. Datskos, “Effect of Temperature on the Formation and Autodestruction of Parent Anions,” *International Journal of Mass Spectrometry and Ion Processes*, Vol. 149 / 150, pp. 59–77, 1995.
 27. S. J. Dale, R. E. Wootton, and A. H. Cookson, “Effects of Particle Contamination in SF₆ CGIT Systems and Methods of Particle Control and Elimination,” in *Gaseous Dielectrics II*, L. G. Christophorou (Ed.), Pergamon Press, New York, pp. 256–265, 1980.
 28. R. E. Wootton, in *Gaseous Dielectrics III*, L. G. Christophorou (Ed.), Pergamon Press, New York, p. 554, 1982.
 29. L. G. Christophorou, I. Sauers, D. R. James, H. Rodrigo, M. O. Pace, J. G. Carter and S. R. Hunter, “Recent Advances in Gaseous Dielectrics at Oak Ridge National Laboratory,” *IEEE Trans. Electr. Insul.*, Vol. EI-19, pp. 550–566, 1984.
 30. D. W. Bouldin, D. R. James, M. O. Pace, and L. G. Christophorou, “A Current Assessment of the Potential of Dielectric Gas Mixtures for Industrial Applications,” in *Gaseous Dielectrics IV*, pp. 204–212, 1984.
 31. D. R. James, M. O. Pace, D. W. Bouldin, and L. G. Christophorou, Oak Ridge National Laboratory Report, ORNL/TM-9017, 1984.
 32. K. B. Miners, M. J. Mastroianni, P. N. Sheldon and D. P. Wilson, “Dew Points of SF₆ / N₂ Mixtures,” in *Gaseous Dielectrics III*, L. G. Christophorou (Ed.), Pergamon Press, New York, pp. 509–516, 1982.
 33. L. G. Christophorou and R. J. Van Brunt, in Ref. [20], pp. 147–175.
 34. M. O. Pace, D. L. McCorkle, and X. Waymel, “Possible High Pressure Nitrogen-Based Insulation for Compressed-Gas-Insulated Cables,” 1995 Annual Report, Conference on Electrical Insulation and Dielectric Phenomena, 95CH35842, pp. 195–198, 1995; “N₂/SF₆ Breakdown in Concentric Cylinders: Consolidation of Data,” Annual Report, Conference on Electrical Insulation and Dielectric Phenomena, San Francisco, October 20–30, pp. 571–576, 1996; M. O. Pace, private communication, 1997.
 35. L. G. Christophorou and S. R. Hunter, in *Electron-Molecule Interactions and their Applications*, L. G. Christophorou (Ed.), Academic Press, New York, Vol. 2, Chap. 5, 1984.
 36. L. G. Christophorou, D. R. James, I. Sauers, M. O. Pace, R. Y. Pai, and A. Fatheddin, “Ternary Gas Dielectrics,” in *Gaseous Dielectrics III*, L. G. Christophorou (Ed.), Pergamon Press, New York, pp. 151–165, 1982.
 37. A. H. Cookson and B. O. Pedersen, “Analysis of the High Voltage Breakdown Results for Mixtures of SF₆ with CO₂, N₂, and Air,” Third Intern. Symposium on High Voltage Engineering, Milan, paper 31.10, 1979.
 38. N. H. Malik and A. H. Qureshi, “A Review of Electrical Breakdown in Mixtures of SF₆ and Other Gases”, *IEEE Trans. Electr. Insul.*, Vol. EI-14, pp. 1–13, 1979.
 39. J. C. Cronin, “Experience in Design and Testing of Gas-Insulated Systems,” in *Gaseous Dielectrics I*, L. G. Christophorou. (Ed.), Oak Ridge National Laboratory Report CONF-780301, May 1978, pp. 116–132.
 40. K. Nakanishi, “New Gaseous Insulation,” *IEEE Trans. Electrical Insulation*, Vol. EI-21, pp. 933–937, 1986.
 41. Y. Qiu and Y. P. Feng, “Investigation of SF₆/N₂,

- SF₆/CO₂, and SF₆/Air as Substitutes for SF₆ Insulation,” Conference Record of the 1996 International Symposium on Electrical Insulation, Montreal, Quebec, Canada, June 16–19, pp. 766–769, 1996.
42. A. H. Cookson, “Electrical Breakdown Studies of SF₆/CO₂/Fluorocarbon Mixtures,” in *Gaseous Dielectrics II*, L. G. Christophorou (Ed.), Pergamon Press, New York, pp.169–178, 1980.
 43. A. Rein and J. Kulsetås, “Impulse Breakdown of SF₆/N₂ Insulation. Influence of Electrode Covering. Polarity Effects,” in *Gaseous Dielectrics III*, L. G. Christophorou (Ed.), Pergamon Press, New York , pp. 315–321, 1982.
 44. A. Fatheddin, M. O. Pace, L. G. Christophorou, and R. Y. Pai, “Impulse Breakdown in SF₆, SF₆/N₂ and SF₆/N₂/Perfluorocarbon Mixtures,” in *Gaseous Dielectrics III*, L. G. Christophorou (Ed.), Pergamon Press, New York, pp. 183–190, 1982.
 45. A. H. Cookson and R. E. Wootton, “Particle Movement and Gas Breakdown in High Pressure Nitrogen and Sulphur Hexafluoride,” Third Intern. Conf. Gas Discharges, London, pp. 385–388, 1974.
 46. S. J. Dale, R. E. Wootton, and A. H. Cookson, “Effect of Particle Contamination in SF₆ CGIT Systems and Methods of Particle Control and Elimination,” in *Gaseous Dielectrics II*, L. G. Christophorou (Ed.), Pergamon Press, New York, pp. 256–265, 1980.
 47. R. E. Wootton, A. H. Cookson, F. T. Emery, and O. Farish, “Investigation of HV Particle-Initiated Breakdown in Gas-Insulated Systems”, Report No. EPRI EL-1007, 1970.
 48. C. M. Cooke, R. E. Wootton, and A. H. Cookson, “Influence of Particles on AC and DC Electrical Performance of Gas-Insulated Systems at Extra-High Voltage,” IEEE Trans. Power App. Syst. Vol. PAS-96, pp. 768–777, 1977.
 49. A. H. Cookson and O. Farish, “Particle-Initiated Breakdown Between Coaxial Electrodes in Compressed SF₆,” IEEE Trans. Power App. Syst., Vol. 92, pp. 871–876, 1973.
 50. A. Diessner and J. G. Trump, “Free Conducting Particles in a Coaxial Compressed-Gas-Insulated System,” IEEE Trans. Power App. Syst., Vol. 89, pp. 1970–1978, 1979.
 51. R. E. Wootton and A. H. Cookson, “AC Particle-Initiated Breakdown in Compressed Gas Mixtures of SF₆ with He, N₂, and CO₂,” 5th International Conference on Gas Discharges, IEE Conference Publication Number 165, pp.177–180, 1978.
 52. M. O. Pace, J. L. Adcock, and L. G. Christophorou, “Particle Contamination in Gas-Insulated Systems: New Control Methods and Optimum SF₆ / N₂ Mixtures,” in *Gaseous Dielectrics IV*, L. G. Christophorou and M. O. Pace (Eds.), Pergamon Press, New York, pp. 377–386, 1984.
 53. J. R. Laghari and A. H. Qureshi, “Flashover Voltages of Cylindrical Insulators in Gas Mixtures,” IEEE Trans. Electr. Insul. EI-16, pp. 373–387, 1981; Sixth Intern. Conf. Gas Discharges and Their Applications, Edinburgh, pp. 240–242, 1980.
 54. M. Eteiba, F. A. M. Rizk, N.-G. Trinh, and C. Vincent, “Influence of a Conducting Particle Attached to an Epoxy Resin Spacer on the Breakdown Voltage of Compressed-Gas Insulation,” in *Gaseous Dielectrics II*, L. G. Christophorou (Ed.), Pergamon Press, New York, pp. 250–254, 1980.
 55. F. Endo (Hitachi, Ltd.) (Private Communication, June 1997).
 56. B. Blankenburg, “Flashover Behavior of Cylindrical Insulators in SF₆, N₂, and SF₆/N₂ Mixtures,” Proc. Third Intern. Symp. on High Voltage Engineering, Milan, paper No. 32.03, 1979.
 57. X. Waymel and C. Boisseau (Electricité de France) (Private Communication, May, 1997).
 58. P. G. Datskos, L. G. Christophorou, and J. G. Carter, “Temperature Dependence of Electron Attachment and Detachment in SF₆ and c-C₄F₆,” J. Chem. Phys., Vol. 99, pp. 8607–8616, 1993.
 59. W. Pfeiffer, V. Zimmer, and P. Zipfl, “Time Lags and Optical Investigations of Pre-Discharge in SF₆ / N₂ Mixtures at Very Fast Transient Voltages,” in *Gaseous Dielectrics VI*, L. G. Christophorou and I. Sauers (Eds.), Plenum Press, New York, pp. 231–236, 1991.
 60. M. Yun and L. Ming, “Investigation of the Toxicological Action of Arc-Decomposed SF₆ and SF₆/N₂,” Sixth International Symposium on High Voltage Engineering, New Orleans, August 28–September 1, paper 32.34, 1989.
 61. Technical Research Report, “Gas-Insulated Transformer II-Utilization,” Electrical Engineering Department, Tsinghua University, Beijing Second Transformer Factory, Beijing, China, 1996 (IDRC Centre File:93-1204-04).
 62. Technical Research Report V, “The Insulation Structure of SF₆ Mixtures GIT and its Test,” Electrical Engineering Department, Tsinghua University, Beijing Second Transformer Factory, Beijing, China, 1995.12 (Project: Gas Insulated Transformer, IDRC Centre File: 93-1204-04).
 63. Technical Research Report I, “Temperature Distribution Test in SF₆ Mixture Gas Insulated Transformer,” Electrical Engineering Department, Tsinghua University, Beijing Second Transformer Factory, Beijing, China, 1995.5 (Project: Gas-Insulated Transformer: II – Utilization, IDRC Centre File:93-1204-04).
 64. E. Kuffel, Private Communication, June 1997.
 65. K. Nakanishi, Private Communication, June 1997.
 66. B. Chervy, H. Riad, A. Gleizes, and J. J. Gonzalez, “Interruption Capability of SF₆/CF₄ and SF₆/C₂F₆ Mixtures,” Eleventh Intern. Conf. Gas Discharges and

- Their Applications, Tokyo, Japan, pp. I-334–I-337, 1995.
67. B. Chervy, H. Riad, and A. Gleizes, “Calculation of the Interruption Capability of SF₆/CF₄ and SF₆/C₂F₆ Mixtures - Part I: Plasma Properties,” IEEE Transactions on Plasma Science, Vol. 24, pp. 198–209, 1996.
 68. B. Chervy, J.-J. Gonzalez, and A. Gleizes, “Calculation of the Interruption Capability of SF₆/CF₄ and SF₆/C₂F₆ Mixtures- Part II: Arc Decay Modeling,” IEEE Transactions on Plasma Science, Vol. 24, pp. 210–217, 1996.
 69. Y. Nakagawa, M. Tsukushi, K. Hirasawa, and Y. Yoshioka, “Interruption Ability of SF₆/N₂ Mixtures and SF₆/CF₄ Mixtures in Puffer Type Gas Blast Circuit Breaker,” Proceedings 4th International Symposium on Switching Arc Phenomena, Łódź, September 22–24, pp. 180–183, 1981.
 70. R. Middleton, V. Koschik, P. Högg, P. Kulkarni and H. Heiermeier, “Development Work for the Application of 245 kV Circuit Breakers Using a SF₆/CF₄ gas mixture on the Manitoba Hydro System,” Canadian Electrical Association Station Equipment Subsection, Electrical Apparatus Section, Engineering and Operating Division, Toronto, Canada, March 1994.
 71. F. M. Clark, *Insulating Materials for Design and Engineering Practice*, John Wiley and Sons, New York, pp. 58–130, 1962.
 72. D. M. Grant, J. F. Perkins, L. C. Campbell, O. E. Ibrahim, and O. Farish, “Comparative Interruption Studies of Gas-Blasted Arcs in SF₆/N₂ and SF₆/He Mixtures,” Proc. 4th Intern. Conf. on Gas Discharges, IEE Conf. Publ. No 143, pp. 48–51, 1976.
 73. R. D. Garzon “The Effects of SF₆/N₂ Mixtures Upon the Recovery Voltage Capability of a Synchronous Interrupter,” IEEE Trans. Power Apparatus and Systems, Vol. PAS-95, pp. 140–144, 1976.
 74. W. M. Leeds, R. E. Friedrich, C. L. Wagner, and T. E. Browne, Jr., C. I. G. R. E., Paris, France, 13–11, 1970
 75. C. E. Sölver, “Thermal Limiting Curves of Full Size Puffer Interrupters. Pure SF₆ and Mixture SF₆/N₂,” Proc. 6th Intern. Conf. Gas Discharges and their Applications, 8–11 September 1980, Heriot-Watt University, Edinburgh, pp. 25–28, 1980.
 76. H. J. Lingal, A. P. Strom and T. E. Browne, “An Investigation of the Arc-Quenching Behavior of Sulphur-Hexafluoride,” Trans. of AIEE, Vol. 72, Part III, pp. 242–246, 1953.
 77. L. D. McConnell and R. D. Garzon, “The Development of a New Synchronous Circuit Breaker,” IEEE Trans. Plasma Science, Vol. PAS-92, pp. 673–681, 1973.
 78. Beehler and L. D. McConnell, “A New Synchronous Circuit Breaker for Machine Protection,” IEEE Trans. Plasma Science, Vol. PAS-92, pp. 668–672, 1973.
 79. H. Naganawa, H. Ohno, M. Iio, and I. Miyachi, “DC Interruption by Spiral Arc in SF₆ / N₂ Mixture,” 8th International Conference on Gas Discharges and Their Applications, Oxford, 1985, pp. 137–140.
 80. A. Lee and L. S. Frost, “Interruption Capability of Gases and Gas Mixtures in a Puffer-Type Interrupter,” IEEE Trans. Plasma Science, Vol. PS-8, pp. 362–367, 1980.
 81. M. Tsukushi, Y. Nakagawa, K. Hirasawa, and Y. Yoshioka, “Application of Theoretical Calculation for Puffer Pressure Rise of Gas Mixtures in Puffer Type Blast Circuit Breakers,” Proceedings 4th International Symposium on Switching Arc Phenomena, Łódź, September. 22–24, pp. 198–202, 1981.
 82. A. Gleizes, M. Razafinimanana, and S. Vacquie, “Temperature Variations in an SF₆/N₂ Mixture Arc Plasma,” Revue Phys. Appl., Vol. 22, pp. 1411–1417, 1987.
 83. A. Gleizes, M. Razafinimanana, and S. Vacquie, “Calculation of Thermodynamic Properties and Transport Coefficients for SF₆/N₂ Mixtures in the Temperature Range 1,000–30,000 K,” Plasma Chemistry and Plasma Processing, Vol 6, pp. 65–78, 1986.
 84. A. Gleizes, I. Sakalis, M. Razafinimanana, and S. Vacquie, “Decay of Wall Stabilized Arcs in SF₆/N₂ Mixtures,” J. Appl. Phys., Vol. 61, pp. 510–518, 1987.
 85. A. Gleizes, A. M. Rahal, H. Delacroix, P. V. Doan, “Study of a Circuit Breaker Arc with Self-Generated Flow: Part I - Energy Transfer in the High-Current Phase,” IEEE Transactions on Plasma Science, Vol. 16, pp. 606–614, 1988.
 86. A. Gleizes, M. Mitiche, P. V. Doan, “Study of a Circuit-Breaker Arc with Self-Generated Flow: Part III - The Post-Arc Phase,” IEEE Transactions on Plasma Science, Vol. 19, pp.12–19, 1991.
 87. A. Gleizes, M. Razafinimanana, and S. Vacquié, “Theoretical and Experimental Study of an Arc Discharge in a SF₆/N₂ Mixture,” 7th International Conference on Gas Discharges and Their Applications, 31 August – 3 September, 1982, London, pp. 85–88, 1982.
 88. H. Sasao, S. Hamano, Y. Ueda, S. Yamaji, and Y. Murai, “Dynamic Behavior of Gas-Blasted Arcs in SF₆-N₂ Mixtures,” IEEE Transactions on Power Apparatus and Systems, Vol. PAS-101, pp. 4024–4029, 1982.
 89. M. C. Siddagangappa and R. J. Van Brunt, “Decomposition Products from Corona in SF₆ / N₂ and SF₆/O₂ Mixtures”, Proc. 8th Int. Conf. on Gas Discharges and Their Applications, Leeds University Press, pp. 247–250, 1985.
 90. R. Probst, DILO Company, Inc. (Private Communication, June 1997).
 91. P. Bolin, Private Communication, June 1997.
 92. L. E. Brothers, Southern Company (Private

- Communication, July 1997).
93. Th. Aschwanden, "Swarm Parameters in SF₆ and SF₆/N₂ Mixtures Determined from a Time-Resolved Discharge Study," in *Gaseous Dielectrics IV*, L. G. Christophorou and M. O. Pace (Eds.), Pergamon Press, New York, pp. 24–33, 1984.
 94. H. C. Doepken, Jr. "Compressed-Gas Insulation in Large Coaxial Systems," IEEE Trans. Power Appar. and Syst., Vol. PAS-88, pp. 364–369, 1969.
 95. C. M. Cooke and R. Velazquez, "The Insulation of Ultra-HV in Coaxial System Using Compressed SF₆ Gas," IEEE Trans. Power App. Syst. **PAS-96**, p. 1491–1497, 1977.
 96. X. Waymel and C. Boisseau, Electricité de France, Private communication, 1997.
 97. P. T. Medeiros, S. R. Naidu, and K. D. Srivastava, "Lighting Impulse Breakdown of Cylindrical Coaxial Gaps in Nitrogen," IEEE Intern. Symp. Electr. Insul., Washington, D. C., 1986, pp. 210–212, 1986.
 98. T. Nitta, Y. Shibuya, Y. Arahata, and H. Kuwahara, "Statistical Approach to the Breakdown Characteristics of Large Scale Gas-Insulated Systems," in *Gaseous Dielectrics*, Oak Ridge National Laboratory Report, CONF-780301, L. G. Christophorou (Ed.), Oak Ridge, TN, pp. 338–354, 1978.
 99. A. Pedersen, in *Gaseous Dielectrics II*, "Evaluation of the Effect of Surface Defects on Breakdown in Strongly Electronegative Gases or Gas Mixtures," L. G. Christophorou (Ed.), Pergamon Press, New York, pp. 201–208, 1980.
 100. J. Berril, J. M. Christensen, and A. Pedersen, "Measurement of the Figure of Merit Related to the Effect of Electrode Surface Defects on Breakdown for Strongly Electronegative Gases or Gas Mixtures," in Seventh Intern. Conf. on Gas Discharges and Their Applications, 31 August–3 September 1982, London, pp. 266–269, 1982.
 101. N. H. Malik, A. H. Qureshi, and G. D. Theophilus, "Static Field Breakdown of SF₆/N₂ Mixtures in Rod/Plane Gaps," IEEE Trans. Electr. Insul., Vol. EI-14, pp. 61–69, 1979.
 102. D. Raghavender and M. S. Naidu, "Lighting Impulse Study of SF₆/N₂ Mixtures With 0.1% to 20% SF₆ content for Rod/Plane Gaps with both Positive and Negative Voltages," 1986 Annual Report, Conference on Electrical Insulation and Dielectric Phenomena, Claymont, Delaware, pp. 327–332, 1986.
 103. Y. Qiu and E. Kuffel, "The Breakdown Strengths of Gas Mixtures Containing a Small Amount of Electronegative Gas in Non-Uniform Field Gaps," Seventh Intern. Conf. on Gas Discharges and Their Applications, 31 August–3 September, 1982, London, pp. 215–218, 1982.
 104. A. Yializis, N. H. Malik, A. H. Qureshi and E. Kuffel, "Impulse Breakdown and Corona Characteristics for Rod/Plane Gaps in Mixtures of SF₆ and Nitrogen with less than 1% of SF₆ Content," IEEE Trans. Power Appar. and Systems, Vol. PAS-98, pp. 1832–1840, 1979; E. Kuffel and A. Yializis, "Impulse Breakdown of Positive and Negative Rod/Plane Gaps in SF₆/N₂ Mixtures," IEEE Trans. Power Appar. Systems, Vol. PAS-97, pp. 2359–2366, 1978.
 105. P. Bolin, Private Communication, August 1997.
 106. R. E. Wootton and S. J. Dale, "60-Hz Breakdown Characteristics of SF₆ and its mixtures with CF₃SF₅, CF₃CFCF₂, and He in Non-uniform Fields," Publication 189, pp. 228–231, 1980.
 107. G. Basile, I. Gallimberti, M. Lissandrin, G. Marchesi, and F. Penon, "Interruption Performances of Circuit Breakers Containing SF₆ Gas Mixtures," IX Intern. Conf. on Gas Discharges and Their Applications, Venice, 19–23 September 1988, pp. 135–138, 1988.
 108. L. Niemeyer, ABB Switzerland, Private Communication, August 1997.
 109. J. Dutton, F. M. Harris, and D. B. Hughes, "Electrical Breakdown of N₂O, SF₆, and N₂O/SF₆ Mixtures," Proceedings IEE, Vol. 121, pp. 223–226, 1974.
 110. K. Nakanishi, D. R. James, H. Rodrigo, and L. G. Christophorou, "The Pressure Dependence of the Dielectric Strength of SO₂ and n-C₄F₁₀," J. Phys. D 17, pp. L73–L76, 1984.
 111. S. R. Hunter and L. G. Christophorou, "Pressure-Dependent Electron Attachment and Breakdown Strengths of Unitary Gases, and Synergism of Binary Gas Mixtures: A Relationship," in *Gaseous Dielectrics IV*, L. G. Christophorou and M. O. Pace (Eds.), Pergamon press, New York, 1984, pp. 115–126.
 112. A. H. Cookson and B. O. Pedersen, "High Voltage Performance of Mixtures of SF₆ with N₂, Air and CO₂ in Compressed-Gas-Insulated Equipment," Fifth International Conference on Gas Discharges, Liverpool, IEE Conference Publication 165, pp. 161–164, 1978.
 113. Y. Qiu and D. M. Xiao, "Dielectric Strength of the SF₆/CO₂ Gas Mixture in Different Electric Fields," Ninth International Symposium on High Voltage Engineering, Graz, Austria, 2255-1–2255-4, 1995.
 114. Y. Qiu, S. Y. Chen, Y. F. Liu, and E. Kuffel, "Comparison of SF₆/N₂ and SF₆/CO₂ Gas Mixtures Based on the Figure-of Merit Concept," 1988 Annual Report, Conference on Electrical Insulation and Dielectric Phenomena, October 16–20, 1988, Ottawa, Canada, pp. 299–304, 1988.
 115. R. E. Wootton, S. J. Dale, and N. J. Zimmerman, "Electric Strength of Some Gases and Gas Mixtures," in *Gaseous Dielectrics II*, L. G. Christophorou (Ed.), Pergamon Press, New York, pp. 137–147, 1980.
 116. J. Berg and E. Kuffel, "Breakdown Voltage Characteristics of SF₆/CF₄ Mixtures in Uniform and Non-Uniform Field Gaps," 1995 Annual Report, Conference on Electrical Insulation and Dielectric

- Phenomena, 95CH35842, pp. 126–129, 1995.
117. M. Akbar and N. H. Malik, “Electrical Breakdown of N_2O/SF_6 , N_2O/CCl_2F_2 , and N_2O/CO_2 Gas Mixtures,” *IEEE Trans. Electric. Insul.*, Vol. EI-20, pp. 581–585, 1985.
 118. Y. A. Safar, N. H. Malik and A. H. Qureshi, “Impulse Breakdown of Negative Rod-Plane Gaps in SF_6/N_2 , SF_6/Air , and SF_6/CO_2 Mixtures,” *IEEE Trans. Electric. Insulation*, Vol. EI-17, pp. 441–450, 1982.
 119. N. H. Malik, A. H. Qureshi, and Y. A. Safar, “DC Voltage Breakdown of SF_6/Air , and SF_6/CO_2 Mixtures in Rod-Plane Gaps,” *IEEE Trans. Electr. Insul.*, Vol. EI-18, pp. 629–636, 1983.
 120. C. C. Chan, M. O. Pace, and L. G. Christophorou, “DC Breakdown Strengths of Some Multicomponent Gas Mixtures in Concentric Cylinder Geometries,” in *Gaseous Dielectrics II*, L. G. Christophorou (Ed.), Pergamon Press, New York, 1980, pp. 149–159, 1980.
 121. Y. Qiu and Y. P. Feng, “Investigation of SF_6/N_2 , SF_6/CO_2 , and SF_6/Air as Substitutes for SF_6 Insulation,” *Conference Record of the 1996 IEEE International Symposium on Electrical Insulation*, Montreal, Quebec, Canada, June 16–19, 1996, pp. 766–769, 1996.
 122. Y. Qiu and D. M. Xiao, “Dielectric Strength of the SF_6/CO_2 Mixture in Different Electric Fields,” *Ninth International Symposium on High Voltage Engineering*, Vol. S2, 1995, paper 2255-1 – 2255-4.
 123. J. Berg and E. Kuffel, “Breakdown Voltage Characteristics of SF_6/CF_4 Mixtures in Uniform and Non-Uniform Field Gaps,” *IEEE 1995 Annual Report: Conference on Electrical Insulation and Dielectric Phenomena*, October 22–25, 1995, Virginia Beach, VA, pp.126–129, 1995.
 124. L. G. Christophorou, D. R. James, I. Sauers, M. O. Pace, R. Y. Pai, and A. Fatheddin, “Ternary Gas Dielectrics,” in *Gaseous Dielectrics III*, L. G. Christophorou (Ed.), Pergamon Press, New York, pp.151–165, 1982.
 125. D. Raghavender and M. S. Naidu, “A Comprehensive Study of Lightning Impulse Breakdown and Cost / Benefit Analysis of Ternary Gas Mixtures Containing Two Electronegative Gases,” *1995 Annual Report, Conference on Electrical Insulation and Dielectric Phenomena*, 95CH35842, pp. 141–144, 1995.

Appendix A

Sales of SF₆ by end-use application [12] (This survey excludes contributions from China and Russia).

YEAR	Utilities and Accelerators	Original Equip. Manufacturers	Magnesium Industry	Electronics Industry	Utilizing SF ₆ Adiabatic Prop.	All Other Uses	TOTAL
(units in metric tons)							
1961	0	91	0	0	0	0	91
1962	14	145	0	0	0	0	159
1963	14	181	0	0	0	9	204
1964	18	204	14	0	0	14	250
1965	32	231	23	0	0	18	304
1966	61	233	27	2	5	23	374*
1967	63	239	32	5	9	27	392*
1968	63	256	36	5	14	32	428*
1969	84	269	41	7	14	36	496*
1970	140	277	44	9	14	44	606*
1971	468	289	71	10	14	72	1,018*
1972	515	263	63	11	14	64	1,017*
1973	700	355	83	11	14	84	1,385*
1974	732	334	46	12	14	72	1,378*
1975	937	559	120	13	14	111	1,754
1976	1,236	760	195	13	14	126	2,344
1977	1,299	924	214	13	14	151	2,615
1978	1,473	1,019	256	13	20	134	2,915
1,979	1,962	1,267	312	15	32	188	3,776
1980	1,861	1,542	321	16	35	216	3,991
1981	2,055	1,397	320	17	54	210	4,053
1982	2,096	1,550	360	18	59	246	4,329
1983	1,966	1,421	374	17	80	233	4,091
1984	2,348	1,859	397	16	108	232	4,960
1985	2,440	1,734	437	20	111	254	4,996
1986	2,717	1,666	431	38	110	300	5,262
1987	2,784	1,641	415	73	146	265	5,324
1988	2,654	1,649	398	100	162	307	5,270

1989	2,832	1,753	363	117	270	323	5,658
1990	3,323	1,722	385	140	220	435	6,225
1991	3,414	2,022	357	151	260	682	6,886
1992	3,312	2,173	266	201	290	645	6,887
1993	3,523	1,847	274	229	290	658	6,821
1994	3,295	2,402	354	262	325	587	7,225
1995	3,126	2,659	399	300	356	617	7,457
1996	3,139	2,795	544	307	344	442	7,571
Subtotal	56,696	39,728	7,972	2,161	3,426	7,857	118,512
1997	3,669	2,155	426	342	472	257	7,321 E
1,998	3,870	2,118	388	378	502	235	7,491 E
1,999	3,711	2,270	362	417	559	212	7,531 E
2,000	3,709	2,281	347	461	584	189	7,571 E
TOTAL	71,655	48,552	9,495	3,759	5,543	8,750	148,426

* For the years 1966-1974, some companies could not provide a breakdown by end use. TOTAL SALES includes an additional 672 metric tons sold in these years. The sum of sales by end use does not equal total sales.

^E Manufacturers provided projections for sales in 1997-2000.

APPENDIX B

Status of Environmental, Economic, and Policy Issues Driving SF₆ Reduction

This appendix is a brief summary primarily of the policy (or regulatory) issues that may drive the need for reduction of SF₆ use and emissions by industry.

Environmental concern is a significant motivation toward improved SF₆ management or use of alternatives. Pure SF₆ has a greater ability to absorb infrared radiation than most other gases and it has a very long atmospheric lifetime. These properties make SF₆ a potent global warming gas. To put this in perspective, the long-term radiative effect of 1 kg of SF₆ emitted exceeds that of 1 kg of CO₂ by a factor of 23,900. This number refers to the direct Global Warming Potential (GWP)₁₀₀, that is, the extra amount of long wavelength radiation absorbed over a 100-year period.

In recent years, the price of SF₆ has increased, thus promoting the possibility of using alternate gases or gas mixtures. An evaluation of the total cost for switching from one gas to another, requires performing a technology impact assessment including a full life-cycle analysis. Among the economic factors to be considered are (i) the price of the compounds and material resources consumed, (ii) the costs of new and modified equipment, (iii) the costs during equipment operation, including energy dissipated, emissions control, repair and servicing, (iv) the cost and effectiveness of recycling, and (v) the cost of additional personnel and training. Other factors to consider include (i) availability and life of the alternative gas, (ii) electrical reliability, certification and testing of equipment, (iii) health hazards and safety, and (iv) equipment size and land use.

Global emission control measures for perfluorocompounds (PFCs) and SF₆ are at present following the general course set for the "major" greenhouse gases in the Framework Convention on Climate Change (FCCC) and the ongoing Conference of the Parties (COP) meetings. In the United Nations' efforts to develop an international protocol (under the FCCC) for stronger controls of greenhouse gas emissions, the European Union (EU) proposed to the Ad Hoc Group on the Berlin Mandate (AGBM) that fluorocarbon control be incorporated into the final treaty. The EU proposal suggested policies and measures that include:

- product standards with respect to leakages of emissions
- use of selected low GWP (global warming potential) instead of high GWP chemical

- reducing emissions through improved equipment and practices in the electrical equipment industry.

The concern over emissions of SF₆ was further addressed in 1996 as the Second Conference of the Parties formally directed countries to inventory SF₆ emissions and report them to the internal governing body. The Intergovernmental Panel on Climate Change has proposed methods for countries to consider for use in inventorying emissions of SF₆ from electric power equipment.

Individual countries are developing proposals to shape stronger climate protocols under development for COP-3 (scheduled for late 1997). The European Union proposal specifically highlights HFCs (hydrofluorocompounds), PFCs (perfluorocompounds), and sulfur hexafluoride in the list of greenhouse gases that countries would be able to control under the protocol. The current intent of the COP-3 meeting is to produce a protocol for greater control of greenhouse gas emissions worldwide.

While several important issues remain unresolved, it appears likely that "the U.S. will propose that negotiations for post-2000 year emissions reductions focus on realistic, binding commitments that will produce real environmental benefits. . . . the United States will continue to seek market-based solutions that are flexible and cost effective. . . . international cooperation of this challenge remains critical to any effective response."²⁰ It is important to recognize that alternatives taking the form of SF₆-containing mixtures or any other PFC gas or mixture will be subject to these evolving regulations.

In the meantime, the U.S. Environmental Protection Agency (EPA) is continuing with the development of a voluntary program for users of SF₆ in electric power equipment. EPA's goal for the program is to reduce U.S. emissions of the gas from equipment through voluntary means. The program is expected to incorporate best work practices, capture and recycling of used gas, and design of business plans for replacement of obsolete equipment with uncontrollable SF₆ leaks. In the long term, EPA desires to work with industry to eliminate emissions. Elimination of emissions will require cooperation from the users of

²⁰Statement to Industry/NGOs on U.S. Intervention at Climate Change Negotiations by T. E. Worth, U.S. Department of State

equipment as well as the manufacturers of such devices.²¹
For further information on the voluntary program, contact Elizabeth Dutrow of EPA at (202) 233-9061 or at <dutrow.elizabeth@epamail.epa.gov>.

²¹Private communication from Elizabeth Dutrow, Environmental Protection Agency, September 29, 1997.

APPENDIX C

Potential Barriers to Using Gas Mixtures as a Gaseous Dielectric

1. Perceived Disadvantages

There are several potential barriers to using even simple mixtures in electrical equipment. These are:

- more difficult gas supply, recovery and recycling procedures,
- monitoring and maintaining proper concentrations,
- challenge of handling leaks and emergencies,
- unknown long-term stability and performance,
- difficult disposal, and
- cost-of-ownership.

The applicability or significance of the perceived disadvantages listed above will depend on the application. It is simple to fill up equipment with a mixture, but for some applications routine maintenance requires the gas to be removed, recycled, refilled, or disposed. This maintenance routine may entail pumping, purging, filtering and separation of the mixture components, replacement of the gases in the desired ratio, and volume percentage analysis. Such a routine must be reliable and rapid, minimizing down time. To accomplish this task will require different chemical management practices and associated technologies from those used for pure SF₆. In particular, this may require more training, equipment development, process automation, and other higher-function gas handling equipment. *However, there seem to be no fundamental limitations to the use of gas mixtures.*

2. Mixed Gas Availability and Costs

It is not viable economically or physically for gas suppliers to provide large quantities of certified mixtures. This is due to the fact that a mixture cannot be significantly compressed and still maintain the appropriate concentrations in the gas phase. Therefore transportation of mixtures would require large tanks of gas mixtures in the gas phase. Similarly, there are limitations associated with certification, storage, and transport of mixtures. Instead, suppliers recommend that gas be delivered certified pure, then stored and mixed on-site as needed [C1].

Major SF₆ producers are in the USA, Italy, Japan, and Germany, see Table C1 [C1, C2]. SF₆ is prepared for

commercial and industrial use in various grades (minimum 99.8 mole percent pure) and delivered in a wide range of volumes and pressures. The most commonly found impurities, include air, nitrogen, water vapor, carbon dioxide, CF₄, SO₂, HF, H₂S, SO₂F₂, SOF₂, COS, and trace metal oxides and metal fluorides. Pure SF₆ is readily available in a wide range of volumes and pressures.

At present, only small quantities of calibrated mixtures of SF₆ with other gases can be purchased from a limited number of gas-suppliers. Dilute mixtures such as 1% to 5% SF₆ in air, helium or nitrogen, and in various grades, are routinely available from most gas-suppliers. Other mixtures are typically treated as specialty orders and prepared in small quantities (single cylinders or lecture bottles of desired partial pressure) to specified tolerance and certification accuracy.

SF₆ is shipped, under the appropriate regulations, in cylinders or in tube trailers in liquid form. The maximum filling density permitted for SF₆ in cylinders is 120 % (i.e. percent water capacity by weight). Nitrogen gas is shipped in cylinders, tube tank cars, and tube trailers according to regulations. Liquid nitrogen is shipped as a cryogenic fluid in vacuum-insulated cylinders, and in insulated portable tanks, tank trucks, and tank cars. Storage standards would be recommended if gas mixtures were to be routinely handled, recovered, recycled or transported.

While there is no limitation to the availability of the gases, cost and policy considerations cannot be overlooked. Costs of servicing are relatively high for compounds that are restricted under national or international regulations. Consequently, the choice of replacements must consider the dynamics of environmental policy (as discussed in Appendix B). An evaluation of the total cost for recover/recycle and switching from one gas to another, requires performing a

TABLE C1. Major producers of sulfur hexafluoride [C1, C2].

Company	Country
Air Products and Chemicals Inc.	USA
AlliedSignal Chemicals	USA
AGA Gas AB	Sweden
Kanto Denka Kogyo Co.	Japan
Asahi Glass Co.	Japan
Ausimont (Montecatini Edison)	Italy
Solvay (Kali-Chemie)	Germany

Source: C. M. A. Nayar, GEC Alsthom, France

technology impact assessment including a full life-cycle analysis. Among the costs to be considered are (1) the compounds and material resources consumed, (2) the new and modified equipment required, (3) equipment operation, including energy dissipated, emissions control, repair and servicing, and (4) additional personnel and training. The cost of using gas mixtures will depend on the relative quality and quantity of the alternative gas required to meet the necessary certification tests and operating specifications. In addition, new equipment would need to be purchased for storage, gas cooling, pumping, leakage testing, mixing, refilling and other servicing practices.

3. Preparing Gas Mixtures and Material Compatibility

Refilling of equipment with mixtures can be time consuming and take considerable time to verify or certify to the accuracy desired. Similarly, to “top-off” equipment that has suffered a loss of pressure, i.e., to restore the original pressure and gas composition, is a greater challenge than for systems containing a single gas. The filling and maintenance of any electrical equipment with gas mixtures will require development of concentration range standards, preparation tolerances, and analytical accuracy specifications. For example, when preparing a mixture of 40%SF₆-60%N₂ an acceptable concentration range of 10% might require a preparation tolerance of 5% of component and a 2% analytical accuracy of component.

Studies examining the chemical and physical properties of SF₆-N₂ mixtures have been partially motivated by an interest in the ability to reduce condensation (effective dew point) of SF₆ in equipment located in cold climates [C3, C4]. The temperature at which SF₆ liquefies depends on the gas pressure. Miners et al. [C3] demonstrated that SF₆-N₂ gas mixtures are non-ideal. From a practical standpoint a manufacturer who uses ideal gas assumptions to predict the mixture dew point temperatures at system-fill pressures of 100 kPa to 500 KPa would under-predict the temperature by as much as 10 °C. These results suggested that similar considerations be given to all SF₆-containing mixtures. In some cases electrical equipment may require designs incorporating heaters to ensure SF₆ fractions remain above the liquefaction temperature (this is, of course, undesirable). But liquefaction is only one among many parameters to be considered when refilling with a selected mixture and re-rating the pressure of the electrical equipment and application.

While SF₆-N₂ mixtures are relatively inert, other mixtures of gases with known stability problems may require special techniques such as passivation or surface

preparation to eliminate the degradation of the unstable component, generation of undesirable byproducts, chemical attack of surfaces or general corrosion. In the case of pure SF₆, considerable progress has been made in improving the reliability of seal and gasket design and protection against environmental conditions, resulting in proven long-term performance. When determining what materials can be used with a gas mixture, it is important that design and manufacture value compatibility with each of the components of the mixture separately as well as all of the components when they are combined. This implies careful choice of material composition, surface finish and contact methods for walls, spacers, inserts, shields, electrodes, O-ring seals, or use of epoxy formulations, glues and resins. Modes of equipment operation and location for use must be considered to design for temperature, pressure, and humidity variations or gradients. For example, equipment with SF₆-N₂ mixtures should be designed for or maintained at temperatures that will prevent water and SF₆ from condensing, thereby losing performance and homogeneity. If a mixture with a condensable component has been subjected to temperatures at or below its saturation temperature, it will need to be re-homogenized prior to the withdrawal of any of the gas. Various techniques have been developed for the latter purpose.

Since SF₆ is packaged as liquefied gas, special precautions need to be observed when filling equipment. This applies to both to a new mixture and the processed gas after recovery and recycling. One concern is particulate matter generated in storage or transfer leading to the recommendation to filter use to protect contamination of the electrical equipment. Another issue is the enhancement of chemical transfer rate by the common practice of warming the gas cylinder or storage vessel. Warming the storage container during transfer minimizes the refrigeration effect caused by the evaporation of SF₆ while allowing its transfer in the gaseous state. If liquid phase transfer of SF₆ is employed, care is required to ensure the gas is completely vaporized before it enters the equipment to avoid over-pressurization or undesirable refrigeration.

4. Mixed Gas Recovery and Recycling

Recovery, recycling, and destruction of SF₆ is possible, such that there is no need for deliberate release into the atmosphere. However, it appears that current practices are such that economical separation of SF₆ from nitrogen is not possible without some venting of SF₆ into the atmosphere. In the event that end-of-life disposal is required, all regulations governing air emissions and waste management should be followed. SF₆ can be

destroyed by thermal decomposition at elevated temperatures (>1100 °C). Such thermal waste treatment furnaces process the sulfur and fluorine constituents to produce naturally occurring materials, e.g. gypsum and fluorspar.

Choice of alternatives should consider the availability of internal treatments to the electrical equipment, including gas absorbent filters, desiccants, and particle filters used in an effort to stabilize the gases and aid recovery and recycling. Gases, whether initially pure or mixed, can be expected to degrade with time due to a variety of factors including contamination caused by moisture and decomposition products. The composition and concentration of potentially toxic by-products is unknown for most mixtures and will be quite variable between applications and equipment.

Lower fluorides of sulfur formed by the decomposition of SF₆ may be removed by gas scrubber/filter systems. Gaseous decomposition products may be absorbed on molecular sieves or on soda lime (50/50 mixture of NaOH and CaO), or on activated alumina (specially dried Al₂O₃). The quantity of decomposition products and the amount of absorbent required to capture all of the products will need to be determined. It has been suggested that a practical rule-of-thumb is to use a weight of absorbent corresponding to 10% of the weight of gas [C5]. The absorbent should be located in the equipment to maximize gas contact, unless both liquid and gas phases are present. In the latter it may be necessary to locate the absorbent in contact with both phases or only the liquid phase. The effectiveness and saturation of absorbents, desiccants and filters will depend on the equipment design, maintenance schedules, temperature, as well as consequence of equipment faults and contamination.

The gas from a faulted breaker, leaking transmission line or transformers, or gas-insulated substation, once treated to remove decomposition products and moisture, may be reused if the material meets device specifications. The key to continued reusability of the gas is to establish purity standards, certification requirements, and recovery / recycling protocols to performance specifications. It takes a combination of factors to achieve this goal:

- Contamination minimization must be built in to electrical equipment design and operation;
- Contamination minimization must be built in to delivery, mixing, recover, and recycle equipment design, operation and chemical management practices; and
- Monitoring of gas condition including electrical properties and chemical properties (e.g., purity, decomposition products, moisture content) must be available; and
- Quality of chemical equipment manufacture, equipment maintenance, and chemical management

practices must be continuously improved.

The desired purpose of gas recycling is to recover the original gas, remove any undesirable byproducts (such as moisture, oil, and particles), verify and possibly correct the mixture composition, and return the gas back to the electrical equipment in a satisfactory certifiable state. A moisture specification of around 30 ppmv (parts-per-million-volume) is typical while the IEC Standard 376 for new SF₆ gas specifies an oil content not to exceed 5 ppmw (parts-per-million-weight). Two international committees (CIGRÉ WG 23-10 TF 01 and IEEE-EI S32; [C6]) are attempting to define purity standards for on-site recycled SF₆. The standards and protocols for recovery and recycling of alternatives could be developed in a similar manner. Draft standard IEEE P1403, which compares air-insulated substations and gas-insulated substations (GIS), mentions that recent advances in GIS construction include sophisticated equipment to reprocess SF₆. Similar integrated technology could be developed for dealing with the potential alternatives.

In Japan, the Task Committee on the Standardization of the Use of SF₆ Gas for Electrical Power Equipment is currently examining the practices for recycling and handling of SF₆ gas. Among the targeted voluntary actions is the reduction of releases of SF₆ at all stages of equipment development, installation, and testing. Targets for recovery are 97% of the purchased gas by the year 2005. This is to be accomplished by the development of economical and large capacity recycling systems which evacuate vessels to higher vacuum. Similar recovery and recycle practices could be implemented for mixtures but have not been explicitly discussed by this task force.

Control of the temperature and pressure is critical to successful reclamation in gas mixtures. In the case of recovering SF₆ from SF₆-N₂ mixtures, the N₂ typically represents a compressible but non-liquefiable component that reduces the overall extraction efficiency, unless higher operating pressures or lower temperatures can be attained. It should be noted that very little thermodynamic data on SF₆-containing mixtures are available in the scientific literature. Computational tools are currently available to help predict some of these missing data [C7]. Efforts to employ these tools may enhance efforts to implement the chemical management of SF₆-containing mixtures as alternative gases. Studies by Mitchel *et al.* [C8] calculated the SF₆ liquid / SF₆ gas / N₂ gas phase equilibrium assuming a constant volume for an initial fill of various blends at several initial pressures at 20 °C, subsequently cooled to -50 °C. They concluded that reclamation of SF₆ from SF₆-N₂ and SF₆-air mixtures is best accomplished by a combination of compression and refrigeration to liquefy the SF₆. Volumetric efficient handling of mixtures is considered to require cooling assisted high-pressure (rather than low-pressure) devices.

Generally, more dilute SF₆ mixtures require lower temperatures and/or higher pressures.

To date, commercially available gas reclamation technology for the electrical industry has been designed primarily for separation, processing, analysis, and compression of nearly pure SF₆ gas. Much of this work has been done by companies specializing in SF₆ processing, working in cooperation with one or more manufacturers of SF₆ insulated equipment. In most cases, gas carts use pressurized liquefaction of SF₆ (via compressors) to minimize the necessary volumetric storage required [C8-C10]. When the stored gas is nearly 100% pure SF₆ this method of reclamation is highly satisfactory and recovery rates greater than 99% yield are possible [C5, C10]. Losses of SF₆ to the environment depend strongly on the SF₆ percentage in the mixture, the operating pressure, the extent of cooling, and the residual pressure remaining in the evacuated volume. Table 2C shows the losses predicted by Probst [C10], using *currently available* technology based on a two-cycle distillation process operating at high pressure (5000 KPa) and low temperature (-40 °C), where liquid SF₆ is withdrawn and the remaining gas cushion is vented, when the purity of liquid gas is to be better than 99%.

Conventional SF₆ gas reclamation carts have limited capability for processing SF₆ containing N₂, air or decomposition byproducts at levels exceeding a few percent [C5, C9]. If gases are heavier than N₂ (for example CF₄) then the SF₆ losses can be substantial. On conventional carts, SF₆ is cycled and liquefied but the nitrogen gas cannot be liquefied. Liquefaction lowers the total pressure in the process tank. Each cycle consists of adding mixed gas until the total pressure equals the initial pressure, followed by additional cooling. If the on-board volumetric storage tank is not sufficiently large, the potential exists for N₂ gas to shut down the compressor at some limiting high pressure. The ultimate capacity of the cart storage is reached when the residual gas is compressed to the maximum safe pressure. At this point the volume being evacuated inside the electrical equipment may still contain some unknown ratio of mixed gas and the storage tank will hold liquefied SF₆ and gaseous N₂. The protocol used to minimize evaporative loss of SF₆ recommends always reaching full capacity before emptying the tank, and when emptying to first transfer the SF₆ liquid and then purge the residual gas [C5, C8].

Practical applications of SF₆-N₂ mixtures where N₂ gas is the predominant gas requires refrigeration to separate and recover the SF₆ efficiently. B. Smith [C5] recommends that in some instances it may be more appropriate to use low pressure-assisted cooling operation instead of high pressure devices [C10]. In this case, pumps are configured to maximize the quantity of gas

TABLE 2C. Estimated losses for recover/recycling procedures

SF ₆ Percentage in Mixture	Expected Losses
> 98%	10
>90%	12
>80%	15
>70%	20
>60%	30
>50%	50

Source: R. Probst, DILO company, Inc. [C10]

withdrawn from the electrical equipment (reaching base pressures on the order of 100 Pa) [C5]. Commercial refrigeration systems are available that use an initial liquefaction of the reclaimed gas (e.g., SF₆ and contaminants), followed by a further liquefaction of the gas phase by sub-cooling of the gas/liquid mixture in a separate column. Nitrogen gas and contaminants can be slowly vented while the entrained SF₆ can be re-liquefied and stored. Continuous sub-cooling of the liquid SF₆ further separates the gases. Once isolated the SF₆ can be continuously recycled to dry and purify the gas.

To assure efficient SF₆-N₂ mixing, the recommended protocol for returning recycled gas to the electrical equipment should be to start with nitrogen gas transfer [C5]. As pressure is equalized between the equipment and SF₆ storage tank, heated SF₆ gas can be transferred from storage tank to electrical equipment until the desired mixing ratio (partial pressure) is obtained. The uniformity of mixing among gas components is important when refilling with recycled gas. This can be accomplished by allowing sufficient time for diffusion, designing equipment with several carefully selected points of gas injection, and by creating turbulence during the mixing period. The rate of recovery varies with process used, for example the recovery can be quite slow (on the order of 10-400 lbs/hr) using conventional gas carts. Such limitations may not exist with refrigeration systems. More complex, low pressure gas carts are typically faster, and recover more gas, then comparable high pressure systems [C5]. Refill and storage does not appear to be a problem. Refilling of any container with or without refrigeration devices or heat exchangers is commercially viable.

If a replacement gas mixture cannot be recovered and recycled in a safe, cost effective, and environmentally protective manner, then no real improvement has been achieved [C11]. Additional study of the chemical and physical properties associated with recovery and recycle of possible replacement gas mixtures needs to be pursued to accelerate the recommendation, testing, and implementation of any alternatives to pure SF₆.

5. Retrofit

A number of manufacturers of electrical equipment and specialty companies have developed methods to retrofit circuit breakers and other devices with vacuum and SF₆ interrupters. Depending on the type of application and equipment it may be reasonable to retrofit equipment with modified gas manifolds, heaters, storage compartments, material coatings, filters and traps, etc. This concept of retrofitting technology has been proven reliable and cost effective in certain specific applications. Past experience in this area can be used to help retrofit devices for use of alternatives.

In some applications, the use of replacement gas mixtures would require considerably higher operating pressures than pure SF₆ requires. Specific equipment designs, construction, and manufacture will have to be evaluated for the ability to accommodate such pressure changes. Otherwise, to maintain similar electrical properties at the same operating pressure, larger and more robust equipment might be required. Larger insulating clearances, improved rupture disks, or whatever the retrofit for pressure requires may only be readily introduced in the design of new equipment. In other cases, the changes in operation may be associated with thermal changes, transport properties, or other mechanisms not readily addressed via retrofitting. The feasibility of electrical and/or thermal derating of existing equipment while purchasing additional equipment will have to be carefully examined. Further research and development into material properties along with gas thermodynamics and kinetics is needed to recommend and implement the retrofit of the installed base of electrical devices.

Any change from the original equipment design, such as substituting a new insulating gas or gas mixture, in existing equipment would require complete resetting and certification of the equipment. Again, there are no fundamental limitations to such testing but there are economic concerns. The testing procedures are described in a number of international and national standards. For example with circuit breakers, the required tests are defined in IEEE Standard C37.09-1979 (Reaff 1988) "Standard Test Procedure for AC High-Voltage Circuit Breakers Rated on a Symmetrical Current Basis." Current practices are such that complete type testing on high voltage electrical equipment can be prohibitively expensive with estimated costs reaching from \$500,000 to \$1,000,000 for each type of breaker tested [C11]. The move to alternate gases would require research, development, and policy changes. These would be intended to provide more cost-effective, rapid and accurate testing and certification procedures.

6. Monitoring and Analysis

Monitoring and analysis are primarily used to determine when maintenance is required and to evaluate the equipment condition and gas quality. This includes monitoring of gas adsorbent column, desiccants, particle filters, and gas scrubbers. Monitoring equipment designed specifically for pure SF₆ applications is currently available and may be useful to monitor SF₆-or alternative gas mixtures. Research and development may be warranted to certify the performance of such equipment with mixtures and advance microprocessor technologies for multiple gas sensing. In all cases, multi-gas testing would measure moisture content and trace contaminants. Because moisture is the most detrimental contaminant in pure SF₆ applications, careful monitoring of humidity will remain an issue with fluorocompound-containing mixtures.

Monitoring equipment for SF₆-containing mixtures or other alternatives must be sensitive to key byproducts and be reliable over long periods of time. In the case of large equipment, such as substations, automated and multipoint sampling would be valuable. To safeguard the environment against leaks from installed and newly manufactured equipment, use of alternatives may require that monitoring systems be developed for installation at transformer and switch-gear stations.

7. References to Appendix C

- C1. H. Withers, Air Products and Chemicals, Allentown, PA, private communication 1997.
- C2. C. Kroeze, *Fluorocarbons and SF₆: Global Emission Inventory and Options for Control*, [Report No. RIVM-773001007] National Institute of Public Health and Environmental Protection (RIVM), Bilthoven, The Netherlands, 1995.
- C3. K. B. Miners, M. J. Mastroianni, P. N. Sheldon, and D. P. Wilson, "Dew Points of SF₆ / N₂ Gas Mixtures," in *Gaseous Dielectrics III* [L. G. Christophorou, Ed., Pergamon Press, NY 1982] pp. 509-516.
- C4. N. G. Trinh and N. Cuk, "Practical Considerations for Industrial Applications of SF₆ / N₂ Mixtures," Canadian Electrical Association Engineering and Operating Div. Trans., Vol. 23, Pt.1, 84-A-60, Canadian Electrical Association, Montreal, 1984
- C5. B. Smith, "Mixed Gas Reclamation," *Cryoquip Technical Bulletin*, Murrieta, CA 1996; *Sulfur Hexafluoride Gas Recycling Handbook* Cryoquip, Murrieta, CA 1997.
- C6. G. Mauthe, L. Niemeyer, B. M. Pryor, R. Probst, H. Brautigam, P. A. O'Connell, K. Pettersson, H.

- D. Morrison, J. Poblitzki, D. Koenig, "SF₆ and the Global Atmosphere," CIGRE Working Group 23.10. "Gas Insulated Substations," *Electra* No. 164, pp. 121-131, 1996; L. Niemeyer, R. Probst, G. Mauthe, H. D. Morrison, J. Poblitzki, P. Bolin, B. M. Pryor, CIGRÉ WG 23-10 Task Force 01, "SF₆ Recycling Guide," 1996.
- C7. D. Friend, NIST, Boulder, personal communication, 1997.
- C8. G. R. Mitchel, J. Castonguay, N. G. Trinh, "Practical Thermodynamics of SF₆ Recovery from SF₆/N₂ or SF₆/Air Mixtures," in *Gas-Insulated Substations* [S. A. Boggs, F.Y. Chu, N. Fujimoto, Eds., Pergamon Press, NY 1985] pp. 437-442.
- C9. R. Probst, *SF₆ Maintenance Equipment*, DILO Company, Inc., Oldsmar, FL.
- C10. R. Probst, "Recycling of SF₆/N₂ Mixtures," DILO Company, Inc., Oldsmar, FL, personal communication, 1997.
- C11. P. Bolin, Mitsubishi Electric Power Products, Inc. and L. Brothers, Southern Company Services, Inc., personal communication 1997.