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NH3 has 4 negative centres of charge/three bonding pairs and one lone pair and BF3 has 3 negative centres of charge/three bonding pairs / OWTTE; (bond

angles) 107° in NH_3 and 120° in BF_3 ; [4] Accept 107.5° for NH_3 . (iv) BF_3 not polar as no net dipole moment / BF bond polarities cancel each other out / symmetrical distribution of charge;

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Advances in Clinical Chemistry

Bread, pasta, noodles ... some of the many ways in which humans consume wheat after processing has taken place. The gluten proteins of wheat grain, which determine the processing properties of wheat flour, have been the subject of intensive study for many years. The structures, genetics and functional properties of this unique group of proteins are the focus of this book. Providing a unique "snapshot" of the most exciting current research in the area, this wide-ranging book encompasses topics such as biotechnology; analysis, purification and characterization; quality testing; and environmental impacts. Contributions come from academia, government laboratories and industry throughout the world, and will be welcomed by practitioners in a variety of fields

including the food, biological and agricultural sciences.

Oligonucleotides represent one of the most significant pharmaceutical breakthroughs in recent years, showing great promise as diagnostic and therapeutic agents for malignant tumors, cardiovascular disease, diabetes, viral infections, and many other degenerative disorders. The Handbook of Analysis of Oligonucleotides and Related Products is an essential reference manual on the practical application of modern and emerging analytical techniques for the analysis of this unique class of compounds. A strong collaboration among thirty leading analytical scientists from around the world, the book provides readers with a comprehensive overview of the most commonly used analytical techniques and their advantages and limitations in assuring the identity, purity, quality, and strength of an oligonucleotide intended for therapeutic use. Topics discussed include: Strategies for enzymatic or chemical degradation of chemically modified oligonucleotides toward mass spectrometric sequencing Purity analysis by chromatographic or electrophoretic methods, including RP-HPLC, AX-HPLC, HILIC, SEC, and CGE Characterization of sequence-related impurities in oligonucleotides by mass spectrometry and chromatography Structure elucidation by spectroscopic methods (IR, NMR, MS) as well as base composition and thermal melt analysis (T_m) Approaches for the accurate determination of molar extinction coefficient of oligonucleotides Accurate determination of assay values Assessment of the overall quality of oligonucleotides, including microbial analysis and determination of residual solvents and heavy metals Strategies for determining the chemical stability of oligonucleotides The use of hybridization techniques for supporting pharmacokinetics and drug metabolism studies in preclinical and clinical development Guidance for the presentation of relevant analytical information towards meeting current regulatory expectations for oligonucleotide therapeutics This resource provides a practical guide for applying state-of-the-art analytical techniques in research, development, and manufacturing settings.

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