**NC DHHS COVID-19 Vaccination Briefing** 

What North Carolina **Practices Need to Know** 

December 15, 2020





NC DHHS COVID - 19 Response

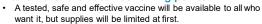
COVID-19 Prevention: Key Messages for December



# **Review & Share the Winter Holidays Guidance**

- · Avoid holiday travel and gatherings with those you don't live with If you must travel or gather: Get tested ahead of time, wear a
- mask all the time, and keep it small and outdoors One-page flyer & detailed guidance (English & Spanish) available at https://covid19.ncdhhs.gov/information/individuals-families-andcommunities/guidelines-get-togethers#winter-holidays

# **Review & Share Vaccines Talking points**



- · The best way to fight COVID-19 is to start first with vaccinations for those most at risk, then reach more people as the vaccine supply increases throughout 2021.
- More information at <a href="https://covid19.ncdhhs.gov/vaccines">https://covid19.ncdhhs.gov/vaccines</a>





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Question during the live webinar



Technical assistance technicalassistanceCOVID19@gmail.com

https://www.communitycarenc.org/newsroom/coronavirus-

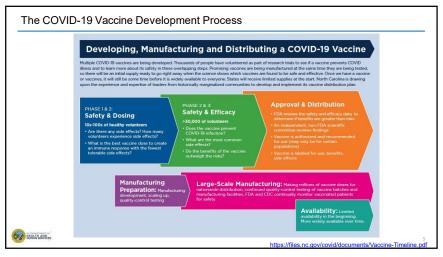
covid-19-information

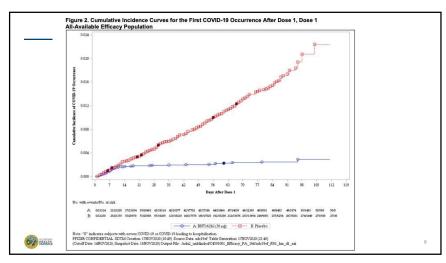
Y'all are doing great... **KEEP CALM CARRY** ON SANT-ANRE

| Agenda       |                                    |  |
|--------------|------------------------------------|--|
|              | Vaccine Authorization and Guidance |  |
|              | NC Vaccine Response Principles     |  |
|              | Priority Groups                    |  |
|              | Overview of Plan                   |  |
|              | Provider Enrollment                |  |
|              | Communications                     |  |
|              | Questions                          |  |
| ISANI SANCES | s                                  |  |

Pfizer Vaccine - Data Brief Phase 3 trial included over 43,000 participants, 42% with diverse backgrounds All others 1% Arizo 4% 16 - 85 years, 46% with co-morbidities (e.g., cancer, heart disease, lung disease, diabetes, obesity, hypertension) 95% effectiveness in preventing illness, 7 days after second dose. • 162/170 cases were in placebo group, 9/10 severe cases were in placebo group Efficacy Data • Uniform effectiveness across age, co-morbidity, demographic groups No waning of protection for at least 2 months after second doses Did not look at data on if a vaccinated person can carry/transmit the virus Applied for EUA 11/20/20, FDA Advisory Committee endorsed 12/10/20 Authorization • Applied for EOA 11/20/20, 10/20 recommendation 12/12/20 Requires ultra-cold storage (-75 degrees Celsius). Storage · Permanent or shipping container refill with dry ice every 5 days up 30 days. 5 days at refrigerated temps 2-dose schedule; 21 days apart (17-21 days), some protection starts 14 days after 1st dose, Dosing Insufficient data to determine protection of 1 dose because almost all got a second dose mRNA technology from the coronavirus's own genes. Tiny piece of genetic material that instructs people's cells Type of make 1 viral protein (spike protein) that triggers immune system to produce antibodies against the COVID Vaccine virus. mRNA technology has been developing for past 2-3 years for other viruses No reports of serious safety during clinical trials. 4 cases of Bell's palsy in vaccine group, same as general rate Safety in population, but will monitor. Temporary reactions (e.g., soreness at site, fatigue, headache, fever) noted 24-48 hours after vaccination, lasts 1-2 days, more after second dose, less with people over 55. Equal percentage of people with and without evidence of prior infection in placebo group became infected (1.3%). "While limited, these data do suggest that previously infected individuals can be at risk of COVID-19 re-infection and could benefit from vaccination.

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# FREQUENCY OF TEMPORARY REACTIONS IN CLINICAL TRIALS BY DOSE AND AGE GROUP, MORE WITH SECOND DOSE, LESS WITH OLDER PEOPLE

| Symptom          | 18-55 year olds |        | > 55 years |        |
|------------------|-----------------|--------|------------|--------|
|                  | Dose 1          | Dose 2 | Dose 1     | Dose 2 |
| Local reaction   |                 |        |            |        |
| Pain at site     | 83%             | 78%    | 71%        | 66%    |
| Redness at site  | 5%              | 6%     | 5%         | 7%     |
| Swelling at site | 6%              | 6%     | 7%         | 8%     |
| Systemic         |                 |        |            |        |
| Fatigue          | 47%             | 59%    | 34%        | 51%    |
| Headache         | 42%             | 52%    | 25%        | 39%    |
| Muscle pain      | 21%             | 37%    | 14%        | 29%    |
| Chills           | 14%             | 35%    | 6%         | 23%    |
| Diarrhea         | 11%             | 10%    | 8%         | 8%     |
| Joint pain       | 11%             | 22%    | 9%         | 19%    |
| Fever            | 3.7%            | 16%    | 1.4%       | 11%    |
| Vomiting         | 1%              | 2%     | 0.5%       | 0.7%   |

3/15,000 people receiving vaccine outside of clinical trial had a severe allergic reaction

# More from FDA Emergency Use Authorization

# Data points from EUA

- Authorized for use for people 16 years of age and older
- Available data on Pfizer-BioNTech COVID-19 Vaccine administered to pregnant women are insufficient to inform vaccine-associated risks in pregnancy.
- Lactation Risk Summary Data are not available to assess the effects of Pfizer-BioNTech COVID-19 Vaccine on the breastfed infant or on milk production/excretion.
- Immunocompromised persons, including individuals receiving immunosuppressant therapy, may have a diminished immune response to the Pfizer-BioNTech COVID-19 Vaccine.
- There is no information on the co-administration of the Pfizer-BioNTech COVID-19 Vaccine with other vaccines

#### Helpful Links

#### Pfizer Website

- Pfizer data briefing document for FDA
- ❖ Full Pfizer-BioNTech COVID-19 Vaccine EUA Letter of Authorization
- Fact Sheet for Healthcare Providers Administering Vaccine (Vaccine Providers)
- \* Fact Sheet for Recipients and Caregivers
- The Advisory Committee on Immunization Practices' Interim Recommendation for Use of Pfizer-BioNTech COVID-19 Vaccine
- ❖ Interim Clinical Considerations for Use of Pfizer-BioNTech COVID-19 Vaccine
- ❖ CDCs COVID-19 Vaccination Communication Toolkit for Medical Center, Clinics, and Clinicians

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#### MORE FROM THE FDA EUA - INGREDIENTS, ALLERGIES

- Ingredients Each 0.3 mL dose of the Pfizer-BioNTech COVID-19 Vaccine contains:
  - 30 mcg of a nucleosidemodified messenger RNA (modRNA) encoding the viral spike (S) glycoprotein of SARS-CoV-2.
  - lipids (0.43 mg (4-hydroxybutyl)azanediyl)bis(hexane-6,1-diyl)bis(2-hexyldecanoate), 0.05 mg 2[(polyethylene glycol)-2000]-N,N-ditetradecylacetamide, 0.09 mg 1,2-distearoyl-sn-glycero-3-phosphocholine, and 0.2 mg cholesterol)
  - 0.01 mg potassium chloride, 0.01 mg monobasic potassium phosphate, 0.36 mg sodium chloride,
     0.07 mg dibasic sodium phosphate dihydrate, and 6 mg sucrose.
- The diluent (0.9% Sodium Chloride Injection) contributes an additional 2.16 mg sodium chloride per dose.
- The Pfizer-BioNTech COVID-19 Vaccine does not contain a preservative.
- Contraindications Do not administer to individuals with known history of a severe allergic reaction (e.g., anaphylaxis) to any component of the Pfizer-BioNTech COVID-19 Vaccine
- Warnings Appropriate medical treatment used to manage immediate allergic reactions must be immediately available in the event an acute anaphylactic reaction occurs following administration of Pfizer-BioNTech COVID-19 Vaccine.

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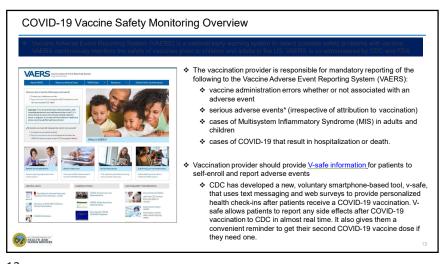
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### MORE FROM FDA EUA - CONSENT

- Due to the FDA Emergency Use Authorization, written informed consent as part of participation in an investigational vaccine development process is no longer required.
- Per the EUA, the vaccination provider, must communicate to the recipient or their caregiver, information consistent with the "Fact Sheet for Recipients and Caregivers" (and provide a copy or direct the individual to the website <a href="www.cvdvaccine.com">www.cvdvaccine.com</a> to obtain the Fact Sheet) prior to the individual receiving Pfizer-BioNTech COVID-19 Vaccine, including:
  - FDA has authorized the emergency use of the Pfizer-BioNTech COVID-19 Vaccine, which is not an FDA-approved vaccine.
  - •The recipient or their caregiver has the option to accept or refuse Pfizer-BioNTech COVID-19 Vaccine.
  - The significant known and potential risks and benefits of Pfizer-BioNTech COVID-19 Vaccine, and the extent to which such risks and benefits are unknown.
  - $\bullet \ \ Information \ about \ available \ alternative \ vaccines \ and \ the \ risks \ and \ benefits \ of \ those \ alternatives.$
- Consent must be obtained prior to vaccination, but that consent can be verbal or written.



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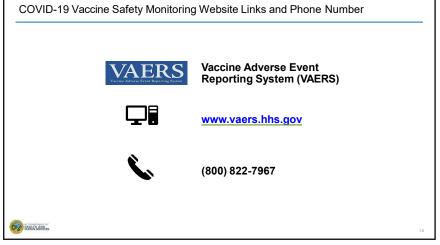
Get vaccinated.
Get your smartphone.
Get started with v-safe.

When your smartphone to tell CDC about a first after getting the CDVID-19 vaccine that uses text messaging and web surveys to provide personalized health check-ins to vaccine recipients following COVID-19 vaccination and serves as an important active surveillance system for adverse events.

V-safe also provides second dose vaccine reminders

All providers who administer COVID-19 vaccine are asked to provide printed hard copies of the v-safe information sheet to each vaccinated individual and counsel them on the importance of enrolling in v-safe.

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CENTERS FOR DISEASE CONTROL AND PREVENTION – ADVISORY COMMITTEE ON IMMUNIZATION PRACTICES (ACIP)

#### · Vaccine Administration

- Pfizer-BioNTech COVID-19 vaccine should be administered alone with minimum interval of 14 days before or after administration with any other vaccines
- Vaccination should be offered to persons regardless of history of prior symptomatic or asymptomatic SARS-CoV-2 infection
- Vaccination should be deferred until recovery from acute illness (if person had symptoms) and criteria have been met to discontinue isolation
- No minimal interval between infection and vaccination however, current evidence suggests reinfection uncommon in the 90 days after initial infection and thus persons with documented acute infection in the preceding 90 days may defer vaccination until the end of this period, if desired

#### · Persons who previously received passive antibody therapy for COVID-19

- Currently no data on safety or efficacy of COVID-19 vaccination in persons who received monoclonal antibodies or convalescent plasma as part of COVID-19 treatment
- Vaccination should be deferred for at least 90 days to avoid interference of the treatment with vaccine-induced immune responses

### • Persons with underlying medical conditions or immunocompromised persons

- Vaccine may be administered to persons with underlying medical conditions who have no contraindications to vaccination
- Persons with HIV infection, other immunocompromising conditions, or who take immunosuppressive medications or therapies might be at increased risk for severe COVID-19 and may still receive COVID-19 vaccine unless otherwise contraindicated



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# **ACIP CLINICAL RECOMMENDATIONS** · Pregnancy and Lactation - COVID-19 and pregnancy - Increased risk of severe illness (ICU admission, mechanical ventilation and death) - Might be an increased risk of adverse pregnancy outcomes, such as preterm birth - mRNA vaccine is not a live virus and the mRNA is degraded quickly by normal cellular processes - If a woman is part of a group (e.g., healthcare personnel) who is recommended to receive a COVID-19 vaccine and is pregnant or lactating, she may choose to be vaccinated. · Contraindications and Precautions - Because of reports of anaphylactic reactions vaccinated outside of clinical trials, - Per EUA Contraindications - Do not administer to individuals with known history of a severe allergic reaction (e.g., anaphylaxis) to any component of the Pfizer-BioNTech COVID-19 Vaccine - ACIP proposed additional guidance: • Persons who have had a severe allergic reaction to any vaccine or injectable therapy (intramuscular, intravenous, or subcutaneous) should not receive the Pfizer-BioNTech vaccine at this time • Vaccine providers should observe patients after vaccination to monitor for the occurrence of immediate adverse reactions: · Persons with a history of anaphylaxis: 30 minutes

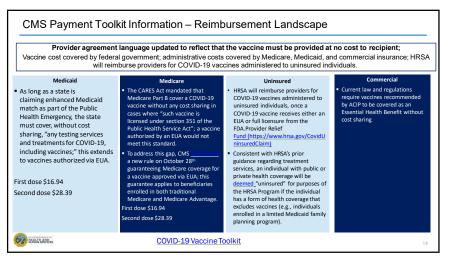
Updates on Remaining Operation Warp Speed Candidates Johnson Johnson NOVAVAX janssen T SANOEL Non-replicating Protein Subunit Protein Subunit 9 ( ) viral vector Phase II/III Phase I/II Phase II/III Est: Early 2021 Est: Early 2021 Est: First half 2021 Est: Early 2021 Doses: 2 (testing halfdose: full-dose regimen v Doses: 1 or 2 (testing both) two full doses) Doses: 1 or 2 (testing Doses: 1 First interim analysis 90% effective with first half-36°F - 46°F 36°F - 46°F 36\*F - 46\*F 36\*F - 46\*F 36°F - 46°F 36°F - 46°F 1B in 2021 1B by mid 2021 2B+ in 2021 Target Supply 100M

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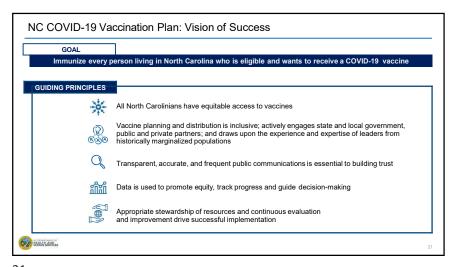
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· All other persons: 15 mins

|                              | Moderna Vaccine  |  |  |
|------------------------------|--|--|--|
| Enrollment                   | Phase 3 trial included 30,000 adult participants 37% with diverse backgrounds. 27% with co-morbidities (e.g., diabetes, heart disease, lung disease, obesity)  |  |  |
| Preliminary<br>Efficacy Data | November 30 Press Release data analysis  94.1% effectiveness in preventing illness, 14 days after second dose.  185/196 cases were in placebo group  30/30 severe cases were in placebo group  Lasts at least 90 days after 2 <sup>nd</sup> dose |  |  |
| Timing of EUA                | Applied for EUA 11/30     FDA Review Dec 17 <sup>th</sup>  |  |  |
| Temperature and<br>Storage   | <ul> <li>Requires storage at -20 degrees Celsius (similar to the chickenpox vaccine) for up to 6 months.</li> <li>Lasts up to 30 days at refrigerated temperatures.</li> </ul>   |  |  |
| Dosing                       | 2-dose schedule     Administered 28 days apart   |  |  |
| Type of Vaccine              | mRNA technology  |  |  |
| Safety                       | No reports of serious safety concerns. Temporary reactions (e.g., fever, soreness at site of injection, fatigue) noted 24-48 hours after vaccination, more after second dose   |  |  |



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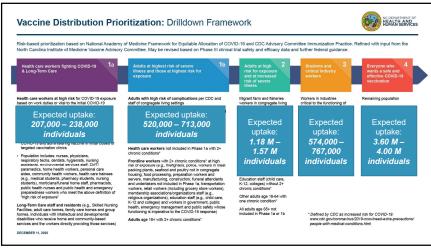
Vaccine Distribution Prioritization: Drilldown Framework NC DEPARTMENT OF HEALTH AND HUMAN SERVICES Risk-based prioritization based on National Academy of Medicine Framework for Equitable Allocation of COVID-19 and CDC Advisory Committee Immunization Practice. Refined with input from the North Carolina Institute of Medicine Vaccine Advisory Committee. May be revised based on Phase III clinical trial safety and efficacy data and further federal guidance. Health care workers at high risk for COVID-19 exposure based on work duties or vital to the initial COVID-19 vaccine response Workers in industries critical to the functioning of society and at increased risk of exposure who are not included in Phase 1 or Phase 2 Adults with high risk of complications per CDC and staff of congregate living settings Migrant farm and fisheries workers in congregate living settings with 2+ chronic conditions\* or age 65+ High risk of exposure is defined as those caring for COVID-19 patients, cleaning areas where COVID-19 patients are admitted, performing procedures at high risk of serosolization (e.g., inhubation, bronchoscopy, suctioning, invasive derital procedures, invasive specimen collection, CPR), handling decelents with COVID-19 and administering vaccine in initial closed or Incarcerated individuals with 2+ chronic conditions\* or age 65+ and jail and prison staff Frontline workers at high or moderate risk of exposure without 2+ chronic conditions Health care workers not included in Phase 1a with 2+ chronic conditions\* Frontline workers with 2+ chronic conditions\* at high risk of exposure (e.g., freelighters, potice, workers in meat packing plants, seafood and poutify not in congregate housing, food processing, preparation workers and senvers, manufacturing, construction, funeral attendants and undertakers not included in Phase 1a, transportation students, pharmacy students, nursing ticians/funeral home staff, pharmacists, and undertakers not included in Phase 1a, transpor workers, retail workers (including grosery store or membership associations/organizations staff (e.g., religious organizations), education staff (e.g., religious organizations), education staff (e.g., religious organizations), education staff (e.g., ker and colleges) and workers in government, pub-health, emergency management and public safety functioning is imperative to the COVID-19 response Other adults age 18-64 with one chronic condition\* Long-Term Care staff and residents (e.g., Skilled Nursing Facilities, adult care homes, family care homes and group homes, individuals with intellectual and developmental disabilities who receive home and community-based \* Defined by CDC as increased risk for COVID-19: www.cdc.gow/coronavirus/2019-ncov/need-extra-people-with-medical-conditions.html Adults age 18+ with 2+ chronic conditions DECEMBER 11, 2020

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#### Advisors

- COVID-19 Vaccine Advisory Committee
  - Purpose: Provide updates from industry and stakeholders to ensure alignment
  - Group of >60 stakeholders
- · Historically Marginalized Populations Advisory Group
  - · Purpose: Identify and address issues related to HMP in the COVID pandemic response
  - Vaccine team presents regularly to HMP Advisory Group for input and partnership opportunities
  - Group of >80 internal and external stakeholders
- COVID-19 Vaccine Communications Advisory Group
  - Purpose: Enhance the development of North Carolina's COVID-19 Vaccine
    Communications Plan and to serve as dissemination partners to extend the reach of the
    communications efforts, especially to prioritized, critical, and historically marginalized
    populations





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# Who is Being Vaccinated in December? Phase 1a

# Where are people in Phase 1a getting vaccinated?

# Health care workers at highest risk for COVID-19 exposure

- · Administering vaccine in initial closed or targeted vaccination clinics
- · In hospitals or local health departments who have received early shipments of vaccine

# Long-Term Care (LTC) staff and residents

- On-site in long-term care facilities in the Pharmacy Partnership for Long-Term Care Program with CVS and Walgreens
- Some will also be vaccinated in local health departments if not with a facility participating in the Pharmacy Partnership program, through other long-term pharmacies, other mobile providers



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# Who is Being Vaccinated in December? Phase 1a

# How will people know if they are in Phase 1a?

# Health care workers at high risk for COVID-19 exposure

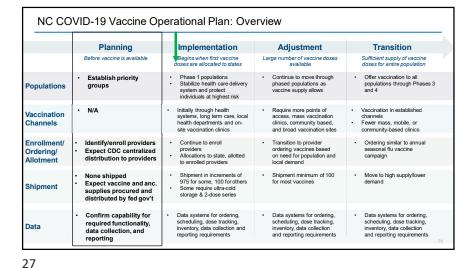
- · Health care employers should determine who meets the criteria for phase 1a
- Health care employers should work with local hospitals or local health departments to determine availability of vaccine and vaccine clinics
- Health care employers should notify employees if they qualify for Phase 1a with instructions for where to be vaccinated

# Long-Term Care (LTC) staff and residents

- · All long-term care staff and residents qualify in Phase 1a
- LTC facilities will be notified when vaccines will be available to be administered to staff and residents.



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Vaccine Journey

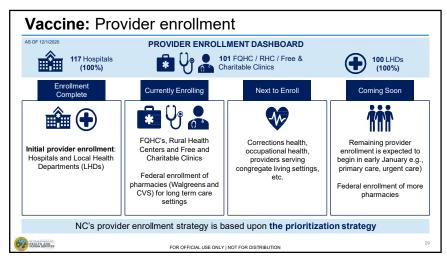
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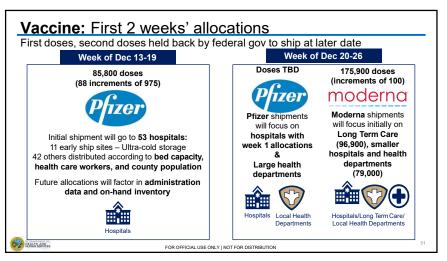
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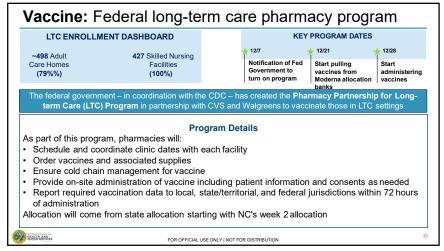
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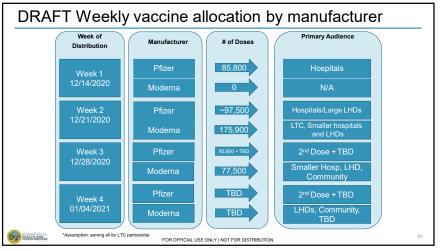
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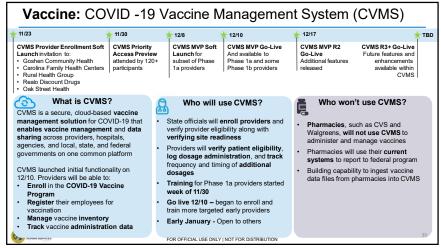
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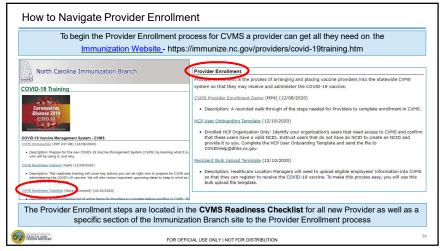
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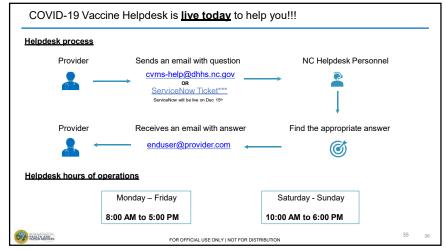
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**CVMS Training and Support Resources** NC DHHS will provide a range of tools and methods for CVMS and vaccine training including; communications, user guides, live trainings, and helpdesk support. Communications: Includes CVMS Live Training: Live training will include Provider Portal announcements, step-by-step demonstrations of key tasks enhancement updates, training event in CVMS, with opportunities to ask invitations, and information about new user ППП questions and do "replays" to take a guides and video demonstrations. closer look with the trainers. A key feature Communications will be tailored to of live training is its high engagement and individual roles and responsibilities interaction from trainees. Helpdesk: email help for all CVMS users during published hours for all CVMS User Guide: Step-by-step guide that related questions. combines text instructions and screenshots to walk users through each task in the CVMS Provider Portal. It breaks down tasks into key steps and includes ServiceNow: CVMS Vaccine Support annotated screen shots and helpfultips. portal will contain a number of Knowledge Articles and FAQ's that will provide information such as self-help. troubleshooting and task resolution. Initial training of Phase 1 enrolled Providers is currently in progress. FOR OFFICIAL USE ONLY | NOT FOR DISTRIBUTION

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#### Questions about CVMS

- How does someone get training on CVMS?/What is the easiest way to get signed up for the CVMS? The quickest and
  easiest way to get training on the CVMS system and all that you need to do to enroll and then use the tool to track vaccine
  and its administration is through the <a href="Immunization Branch Website">Immunization Branch Website</a>. All appropriate materials, check lists for enrolling, and
  then the steps to complete once enrolled are contained within. There is also links to training documents and recordings for
  your end user learning.
- Will the CVMS have a guided questionnaire/logic to help clinicians decide what phase people of distribution that
  their patients fall into? CVMS will automatically determine Priority Tier and Eligibility for recipients in a future release, so
  health care providers will only have to confirm. The Readiness Checklist contains a summary of the prioritization approach
  that North Caroline is currently following.
- Will CVMS integrate into EHRs, including CureMD and Patagonia, which covers the majority of health departments in the state? CVMS does not currently integrate with any electronic health record systems. This is an area that the State is investigating for future enhancements for CVMS to help reduce the amount of double entry of data and to streamline the Healthcare Providers' experience
- How will CVMS works with NCIR and how will the medical home know that a patient got a vaccine from a pharmacist CVMS will interface with NCIR to store vaccine info? The state is using CVMS to track all COVID-19 vaccines administrated across the State. CVMS will interface with NCIR to capture complete immunization information. The State is exploring how to integrate the COVID-19 vaccine administration data from pharmacies participating in the federal Pharmacy Partnership for Long-Term Care Program into CVMS.



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#### **COVID-19 Vaccine Communications**

# **North Carolina's Commitment**

#### Create a proactive, inclusive, evidence-based communications plan that:

- Is guided by research in understanding barriers, values, and motivations for vaccine uptake across different populations
- · Leads with transparency with early and frequent communication about process and plans
- Determines proactive and culturally sensitive and linguistically responsive communication approaches for critical populations as well as the general public
- Communicates clearly and in an impactful way with all stakeholders from start to finish in appropriate languages with tailored and tested messages for target populations
- · Engages trusted community leaders and sources to promote trust



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### **COVID-19 Vaccine Communications**

# **North Carolina's Commitment**

Provide early, transparent, consistent, and frequent communications so that North Carolinians:



Trust the information that they receive from NC DHHS and local health departments about COVID-

19 vaccinations



Understand the benefits and risks of COVID-19 vaccinations



Make informed decisions about COVID-19 vaccinations



Know how and where to get a COVID-19 vaccination

# ISAMILANS:

NC's Communication Strategy Informed by Research

# Addressing Vaccine Confidence – Actionable Data

Many North Carolinians are hesitant about COVID-19 vaccines, particularly Black/African American populations due to longstanding and continuing racial injustices in our health care system.

North Carolina didn't need another poll to tell us people had concerns. Instead, we partnered with the Neimand Collaborative and Artemis Strategy Group to uncover the underlying drivers of awareness, choice and action in health care decisions – **actionable data**.

#### Our research:

- Measures experience, attitudes, knowledge/familiarity and potential barriers with health information and vaccines broadly, and COVID-19 specifically, to identify:
  - · Perceived benefits and risks of COVID-19 vaccinations;
  - Emotional motivations for and against COVID-19 vaccination; and,
  - Trusted sources and spokespeople about COVID-19 vaccinations.



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# NC's Communication Strategy Informed by Research

# Methodology

Surveys conducted from November 10 - November 22, 2020

· Online survey of 1922 North Carolina residents aligned to census data

| North Carolina Sub-Population           | Number of People Who Completed Surveys |
|---|--|
| Rural County Residents                  | 748                                    |
| Urban County Residents                  | 667                                    |
| Suburban/Regional City County Residents | 490                                    |
| COVID Critical County Residents         | 315                                    |
| Blacks/African Americans                | 441                                    |
| Hispanics/Latinx                        | 180                                    |
| American Indians                        | 40                                     |
| Health Care Workers                     | 119                                    |

#### Qualitative interviews conducted from November 28 - December 8, 2020

- 30 in-depth interviews were conducted via Zoom or phone with Black/African American,
- Hispanic/LatinX and American Indian North Carolina adult respondents. Mix of rural, urban, suburban
- · About one third of them were Health Care Workers



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# NC's Communication Strategy Informed by Research

# **Summary Findings and Campaign Implications**

Potential early adoption is weak. Less than half of North Carolina residents are both adherent health decision makers (they tend to follow their doctor's recommendations) and see greater reward than risk in a vaccine—yet a significant number express hesitancy.

The COVID vaccine is not a normal vaccination product. It's new and perceptions of and experiences with other vaccines don't necessarily apply.

Most people are taking a wait and see approach, regardless of demographics. Across demographics, women are the most hesitant—they want to make the right decision for their families.

Hesitancy is driven by legitimate concerns about testing, safety, side effects, effectiveness, "warp speed," and political polarization. These concerns must be addressed *before* any discussion of potential benefits, which are clear to the majority of North Carolinians.

The messengers are 90% of message effectiveness. There is less nuance in messaging than there is messengers. The top three most compelling messages were the same across race and ethnicity. Public health officials are respected, but people also need to see the positive experiences of peers and community leaders.

Vaccine supply and vaccination experience play a large role in communications among a public eager for a cure but waiting to see the positive experiences of "people like them" and a diverse range of others.

COVID-19 Vaccine Message Strategy

- Don't frighten people into wanting to take the shot. They already fear & take COVID seriously. Acknowledge vaccine fears and hesitancy as valid
- · Give people honest information about vaccine development, testing, safety, reactions.
- Build trust in and during the prioritized vaccine rollout: Confidence to frontline workers, patience to eager early adopters, and witness to those who are waiting and seeing.
- · Direct people to "their spot" for reliable information: Official sources or community/peers
- Solve for the logistics of getting people to vaccination sites that may not be connected to their everyday health experiences and health care.
- · Assure everyone of equitable and inclusive access.
- · Have a clear call to action that works across all campaign phases and compliments the 3W's



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#### **COVID-19 Vaccine Message Themes**



Convey Safety in Development Process

Great care has been taken to make sure COVID-19 vaccines are safe and effective.

Scientists had a head start. The vaccines were developed quickly, they were built upon years of work in developing vaccines for similar viruses.

Testing was thorough and successful. More than 70,000 people participated in clinical trials for two leading vaccines to see if they are safe and effective. To date, the vaccines are 95% effective in preventing COVID-19.



Demonstrate Commitment to Transparency & Inclusivity

North Carolina is drawing upon the experience and expertise of leaders from historically marginalized communities to develop and implement its vaccine plan.



Set Expectations

Those who need it most will get it first. A tested, safe and effective vaccine will be available to all who want it, but supplies will be limited at first. The best way to fight COVID-19 is to start first with vaccinations for those most at risk, then reach more people as the vaccine supply increases throughout 2021.



Make the Call to Action

You have a spot. Take your shot. Continue to practice the 3W's until everyone has their shot at fighting COVID-19







